

# DIRECTORY OF SERVICES



**Washington State  
Department of Health**

## **Public Health Laboratories**

**July 2006**

1610 NE 150th Street  
Shoreline, WA 98155-7224  
(206) 418-5400 FAX (206) 418-5445

# **PUBLIC HEALTH LABORATORIES**

## **DIRECTORY OF SERVICES**

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# DOH KEY PERSONNEL PHONE LIST

## PUBLIC HEALTH LABORATORIES:

Website: <http://www.doh.wa.gov/phl/default.htm>

### Director, Public Health Laboratories:

*Romesh Gautom, Ph.D.* .....(206) 418-5450  
[Romesh.Gautom@doh.wa.gov](mailto:Romesh.Gautom@doh.wa.gov)

### Administration:

Laboratory Main Office .....(206) 418-5400  
Main FAX .....(206) 418-5445

### Laboratory Office Directors:

Environmental Laboratory Sciences  
*Nathan Lacy, PhD* .....(206) 418-5520  
Email: [Nathan.Lacy@doh.wa.gov](mailto:Nathan.Lacy@doh.wa.gov)  
Laboratory Operations and Technical Support  
*Lain Knowles* .....(206) 418-5490  
Email: [Lain.Knowles@doh.wa.gov](mailto:Lain.Knowles@doh.wa.gov)  
Newborn Screening  
*Mike Glass* .....(206) 418-5470  
Email: [Mike.Glass@doh.wa.gov](mailto:Mike.Glass@doh.wa.gov)  
Public Health Microbiology  
*Jinxin Hu, PhD* .....(206) 418-5460  
Email: [Jinxin.Hu@doh.wa.gov](mailto:Jinxin.Hu@doh.wa.gov)

### Business Offices:

Central Receiving.....(206) 418-5413  
Mail Room .....(206) 418-5579  
Procurement .....(206) 418-5412

### Public Health Laboratories Safety and Quality Assurance Program

QA and Safety Officer  
*Steve LaCroix,* .....(206) 418-5437  
Email: [Steve.Lacroix@doh.wa.gov](mailto:Steve.Lacroix@doh.wa.gov)  
FAX.....(206) 418-5485

## Public Health Laboratories Training Program

Website: <http://www.doh.wa.gov/EHSPHL/PHL/train.htm>

### Program Manager

*Shelley Lankford*, .....(206) 418-5401

Email: [Shelley.Lankford@doh.wa.gov](mailto:Shelley.Lankford@doh.wa.gov)

Administrative Assistant .....(206) 418-5402

FAX .....(206) 418-5445

## PHL Laboratory Units

Bioterrorism Response .....(206) 418-5471

Chemical Incident Response .....(206) 418-5643

### Drinking Water

Bacteriology .....(206) 418-5489

Chemistry .....(206) 418-5492

MPA .....(206) 418-5489

Enteric Bacteriology .....(206) 418-5456

Food Bacteriology .....(206) 418-5442

Mycobacteriology (TB) .....(206) 418-5473

### Molecular Microbiology

PFGE .....(206) 418-5561

PCR .....(206) 418-5462

Parasitology .....(206) 418-5469

Radiation Chemistry .....(206) 418-5486

### Shellfish

Biotoxins .....(206) 418-5443

Marine Water Bacteriology .....(206) 418-5443

Food Microbiology .....(206) 418-5442

Serology .....(206) 418-5222

### Sexually Transmitted Diseases

Chlamydia .....(206) 418-5607

Gonorrhea (GC) .....(206) 418-5453

HIV/AIDS .....(206) 418-5457

Syphilis .....(206) 418-5622

Special Pathogens Surveillance (Reference) .....(206) 418-5452

Virology .....(206) 418-5458

## LABORATORY QUALITY ASSURANCE

Website: <http://www.doh.wa.gov/hsqa/fsl/LQA/Home.htm>

### Program Manager

*Susan Walker* .....(206) 418-5418

Email: [Susan.Walker@doh.wa.gov](mailto:Susan.Walker@doh.wa.gov)

Administrative Assistant .....(206) 418-5600

FAX .....(206) 418-5505

## NON-LABORATORY PROGRAMS/FUNCTIONS LOCATED AT THE PUBLIC HEALTH LABORATORIES

### Office of Epidemiology-Communicable Disease Epidemiology (CD Epidemiology)

Senior Epidemiologist

*Jo Hofmann, MD*.....(206) 418-5510

Email: [Jo.Hofmann@doh.wa.gov](mailto:Jo.Hofmann@doh.wa.gov)

Administrative Assistant (24-hour number) .....(206) 418-5500

FAX .....(206) 418-5515

### Divisional Emergency Response Planning Advisor

*Robert Arthur Soldier* .....(206) 418-5542

Email: [Robert.Soldier@doh.wa.gov](mailto:Robert.Soldier@doh.wa.gov)

FAX.....(206) 418-5445

### Washington Electronic Disease Surveillance System

*Michael Davisson*.....(206) 418-5420

Email: [Michael.Davisson@doh.wa.gov](mailto:Michael.Davisson@doh.wa.gov)

## OTHER FREQUENTLY CALLED NUMBERS

AIDS Hot Line.....1-800-272-AIDS

Drinking Water Hot Line .....1-800-521-0323

FDA Seafood Hot Line .....1-800-FDA-4010

PSP/Domoic Acid 24-hour Information Line .....1-800-562-5632

Washington State Consumer Assistance Line .....(206) 753-2870

TOLL FREE.....1-800-525-0127

Washington State Basic Health Plan – Insurance .....1-800-773-9872

# PUBLIC HEALTH LABORATORIES



## GENERAL INFORMATION

### MISSION STATEMENTS

**Department of Health Mission:** *The Department of Health Works to Protect and Improve the Health of People in Washington State.*

**Public Health Laboratories Mission:** *Provide a wide range of diagnostic and analytical functions for the assessment and surveillance of infectious/communicable, heritable/genetic and chronic diseases as well as environmental contamination. Improve the quality assurance and analytical performance of clinical and environmental laboratories through training and consultation as well as providing scientific and managerial leadership in developing public health policy.*

### PUBLIC HEALTH LABORATORIES OVERVIEW

#### History

The Washington State Public Health Laboratories (PHL) were established by the legislature in the early 1900's. The laboratories were first located in downtown Seattle in the Alaska Building. The Public Health Laboratories were later moved to the Smith Tower Building and remained there until 1985. In 1982, work was begun on a new facility located a just north of Seattle in the City of Shoreline. The PHL took up residence in the new building in 1985. The laboratories are named in honor of W.R. Giedt, who was the director of the PHL during the period of its greatest changes and growth from 1943 to 1971. Under his leadership the PHL met significant challenges in clinical and environmental public health, and adopted new technologies as soon as they were proven reliable.

Over the past ten years the PHL has focused on the development and implementation of new technologies to provide scientific support focused on improving public health at local, state and



national levels. Dr. Gautom has been instrumental in developing a PFGE procedure that produces results within 24 hours for a variety of pathogens (e.g. E. coli O157:H7, Salmonella, Shigella, etc.) and became the backbone for the national PulseNet system, operated by the CDC, to track national foodborne disease outbreaks.

## **Our Clients**

Primary users of the laboratories include preventive medicine programs at the state, county and federal level; hospitals; public health and medical laboratories seeking reference or consultation services; laboratories desiring certification; other agencies desiring public health laboratory services; and physicians seeking assistance in diagnosing rare or unusual diseases (botulism, rabies, diphtheria, etc.). In addition, programs and agencies concerned with environmental problems make extensive use of the laboratories.

## **Laboratory Services**

The laboratories are engaged in activities designed to aid in the diagnosis, treatment, and prevention of communicable, chronic, congenital and genetic diseases; to assess the general health of the population; to help safeguard a healthful environment; and to assure high quality work within the health and environmental laboratory community. The laboratories provide diagnostic and follow-up services in the areas of newborn screening, food poisoning, surveillance studies of etiologic agents in the areas of bacteriology, virology, serology and parasitology, radiation chemistry, pesticide residue analysis, and many other disciplines. Training, certification and consultation activities are also provided by the State Public Health Laboratories.

As the state's reference clinical laboratory, the PHL provides local health departments, hospitals, clinics and commercial laboratories with a wide range of services including identification and confirmation of unknown pathogenic organisms, consultation on laboratory methodology and training in current laboratory issues and techniques. As a provider of services to local, state and federal agencies, the PHL is often the focal point for coordinating investigations and mediating the transfer of information between agencies. The staff at the PHL test clinical and environmental specimens/samples associated with known and potential disease outbreaks and work with Epidemiology, nursing and environmental health staff to identify possible sources for outbreaks. The PHL staff performs, on an average, 753,800 tests each year for sexually transmitted diseases, foodborne diseases, virus isolation and viral serology, mycobacteriology, environmental microbiology, enterics, parasitology, microbial identification, biotoxins, metals, inorganic chemistry, biotinidase deficiency, congenital adrenal hyperplasia, congenital hypothyroidism, galactosemia, PKU (phenylketonuria), and sickle cell disease in newborns.

## **Response To Bioterrorism and Chemical Terrorism**

DOH (PHL) is participating in a national network called the Laboratory Response Network (LRN) initiated by the Centers for Disease Control and Prevention, Atlanta. The LRN is a collaborative approach between public and private laboratories and is focused heavily on improving laboratory-based bioterrorism and chemical terrorism response capabilities in the United States. Hospital and private laboratories are most likely to be first to receive patient specimens containing etiological agents used in a covert act of bioterrorism and laboratory professionals must be trained to identify microbial pathogens likely to be used for bioterrorism. Laboratorians must know how to safely collect, transport, and process specimens containing

biological agents associated with bio-threat acts and specimens to be analyzed following chemical-threat attacks.

### **The PulseNet Foodborne Disease Surveillance System**

The Centers for Disease Control and Prevention (CDC) in Atlanta, GA, in a cooperative effort with state/local public health agencies, other federal agencies and specialists in the private sector, has developed a foodborne surveillance monitoring system known as PulseNet. PulseNet is an early warning system that allows participating public health laboratories to share critical foodborne disease surveillance information, effectively reducing the time needed to respond to regional and national outbreaks of foodborne disease. Using PulseNet, Pulsed-Field Gel Electrophoresis (PFGE) images and essential demographic information are shared between experts in the investigation of foodborne disease. Bacterial strains, such as *E. coli* O157:H7 and *Salmonella* that may be causing a local foodborne outbreak in one part of the country can be quickly compared with isolates from another locale helping to identify problems where a food source is causing a larger outbreak than first recognized. The PulseNet server is connected to the Internet and is accessible to selected states participating in the PFGE Project, allowing test results to be transmitted quickly and easily between laboratory sites.

### **Outbreak Response**

During 1996-1997 the Microbiology section began developing advanced molecular biology testing capabilities for bacterial and viral pathogens. The methodologies have allowed the PHL to improve the testing services offered to its customers and also to initiate new research projects. Since 1997 the PHL has been testing samples (nasopharyngeal swabs) submitted for *Bordetella pertussis* by PCR. The PHL has conducted two extensive and divided studies with UW to compare various methodologies for detecting *B. pertussis* from clinical samples.

The Public Health Microbiology staff has been directly involved in the investigation of sporadic cases and outbreaks related to *Escherichia coli* O157:H7, *Salmonella*, *Shigella*, *Campylobacter*, *Vibrio parahaemolyticus*, Enterotoxigenic *E. coli*, Methicillin-resistant *Staphylococcus aureus*, Vancomycin-resistant *Enterococcus*, Norwalk-like virus, rubeola, rubella and influenza, to name a few. The team approach of microbiology and epidemiology staff has led to timely intervention for outbreak investigations. For example, during the months of July and August 1999, a unique cluster of 35 cases of *E. coli* O157:H7 was recognized through routine PFGE surveillance testing at the PHL. Patients linked to the cluster reported swimming in a shallow sectioned-off area of a popular swimming lake in southwest Washington. Microbiologists from our Environmental Section were able to isolate *E. coli* O157:H7 from sediment samples collected from the Battleground Lake. This was the first documented report isolating *E. coli* O157:H7 from lake sediment. Subsequently, the PFGE profile from the sediment isolate was found to be identical with all 35 human *E. coli* O157:H7 cases. The PulseNet system helped to provide assurances that the outbreak was not a large multi-state problem but one localized in southwest Washington.

### **PHL Organization**

The Washington State Department of Health is comprised of six divisions. The W.R. Giedt Public Health Laboratories belong to the Division of Epidemiology, Health Statistics and Public Health Laboratories, otherwise known as EHSPHL. The Public Health Laboratories (PHL) are

physically located approximately 10 miles north of downtown Seattle in the city of Shoreline, Washington. The PHL are divided into four major offices, each of which report to the Laboratory Director who in turn reports to the Assistant Secretary for the EHSPHL Division. The offices that comprise the PHL are the Office of Environmental Laboratory Sciences, the Office of Newborn Screening, the Office of Public Health Microbiology and the Office of Laboratory Operations.

#### **Office of Environmental Laboratory Sciences**

The Office of Environmental Laboratory Sciences has approximately thirty technical staff members and is divided into two main sections: Environmental Microbiology and Environmental Chemistry and Radiation. This office is comprised of 7 units that include the Radiation Laboratory, Chemistry Laboratory, Water Microbiology Laboratory, Food Laboratory, Biotoxins Laboratory, Chemical Terrorism Response and Parasitology Laboratory. These laboratory units provide a wide variety of testing of environmental samples and clinical specimens and are certified by several federal programs that include the EPA, FDA, College of American Pathologists and the Nuclear Regulatory Commission.

#### **Office of Laboratory Operations and Technical Support**

The Office of Laboratory Operations and Technical Support has approximately twenty technical and administrative staff. This office is divided into two sections, Administration and Operations and Technical Support. The administration is a group of staff that provides support for the PHL offices and programs. The Operations and Technical Support group provides building maintenance, fiscal management, mailroom and receiving functions, security, The Safety And Quality Assurance Program and The Training Program.

#### **Office of Newborn Screening**

The Office of Newborn Screening has approximately twenty-five technical, follow-up and support staff. The Newborn Screening Program tests every infant born in Washington to detect and prevent the developmental impairments and life-threatening illness associated with phenylketonuria (PKU), congenital hypothyroidism, congenital adrenal hyperplasia, biotinidase deficiency, galactosemia and sickle cell disease (and other clinically significant hemoglobinopathies). The program provides appropriate follow-up and referral of those that screen positive to assure prompt diagnostic and treatment services. In addition, the program provides long-term tracking of affected children to assure continued access to appropriate comprehensive health care. The Office of Newborn Screening screens 150,000 specimens and conducts 670,000 tests every year.

#### **Office Of Public Health Microbiology**

The Office of Public Health Microbiology has approximately thirty technical and support staff. Reference capabilities in this office include diagnostic and surveillance services that focus on foodborne disease, sexually transmitted diseases, virus isolation and viral serology, mycobacteriology, and a variety of new forms of molecular technology. Individual units within the laboratory are

headed by leading experts in the field who work together with the Office of Epidemiology, housed in the same facility, on a daily basis. Virology, serology, HIV, and Chlamydia laboratories perform a variety of conventional, serological and molecular tests to rapidly identify disease agents and characterize viral and bacterial pathogens. Standard tests performed by these laboratories include influenza, rabies, syphilis, EIA and western blot for HIV, Gen-Probe Optima II Combo for Chlamydia/GC, IgG and IgM for rubeola. This office also has a state of the art molecular diagnostics unit that uses DNA based technologies including pulsed field gel electrophoresis (PFGE) and Polymerase chain reaction (PCR) to assist the Office of Epidemiology with outbreak investigations.

## **Using the Directory of Services**

The Directory of Services has been prepared to aid the user in properly utilizing the laboratories services. Information is presented on what is available, how to use it and whom to contact. The Directory contains the telephone numbers of persons responsible for the various disciplines within the PHL. In the interest of providing timely service, users are encouraged to call the laboratory unit to address specific questions. For meaningful results in all areas, an appropriate sample, properly collected and transported, along with adequate identifying information is necessary. Turn-around times are measured in working days. Fees, if applicable, are noted in the directory (all fees are subject to change).



# HOW TO REACH THE PUBLIC HEALTH LABORATORIES



## 24-HOUR EMERGENCY TELEPHONE SERVICE

**(206) 418-5500**

Dialing this phone number will connect the caller to the emergency contact phone operated by the Communicable Disease Epidemiology staff. The person who answers the phone will contact the appropriate laboratory staff.

Hours: 8 a.m. to 5 p.m. on weekdays. Closed on weekends and state holidays, which include New Year's Day, Martin Luther King Jr. Day, Presidents Day, Memorial Day, Independence Day, Labor Day, Veterans Day, Thanksgiving, the day after Thanksgiving and Christmas Day.

**Laboratory Street Address:** 1610 N.E. 150th Street  
Shoreline, Washington 98155

**Mailing Address:** Washington State Public Health Laboratories  
PO Box 550501  
Shoreline, Washington 98155-9701

**Parking:** Free parking is available on-site

### Driving Directions to Laboratory:



#### **I-5, Northbound**

Take N.E. 145th Street exit (Exit #175). After exiting, move to the far right lane. Turn right at traffic light onto N.E. 145th Street (eastbound). Proceed in the left lane on 145th to the next traffic light at 15th Ave. N.E. Turn left onto 15th Ave. N.E. Travel four blocks on 15th Ave. N.E. (northbound) to N.E. 150th St. Turn right onto N.E. 150th. After two blocks, the State Laboratories are on the left.



#### **I-5, Southbound**

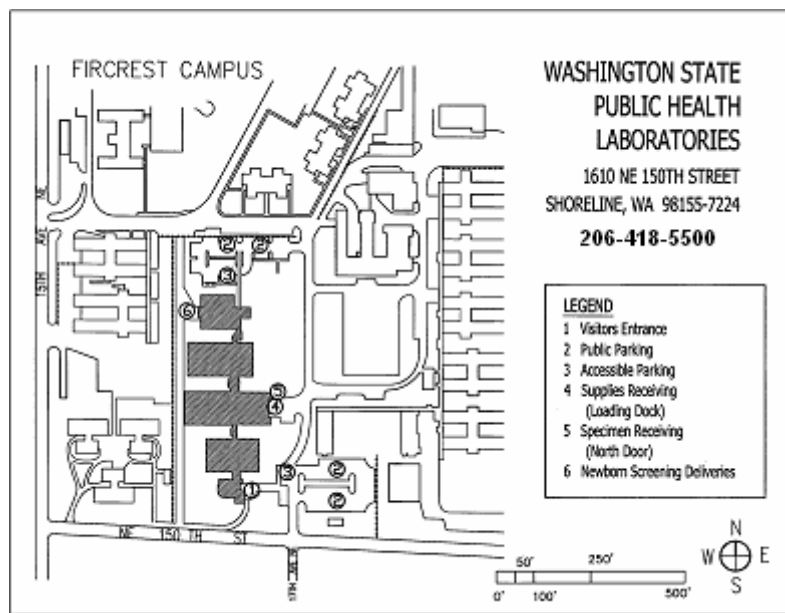
Take N.E. 145th Street exit (Exit #175). After exiting, stay in the left lane of the off ramp. Turn left at traffic light onto N.E. 145th Street (eastbound). Proceed in the left lane on 145th to the second traffic light at 15th Ave. N.E. Turn left onto 15th Ave. N.E. Travel four blocks on 15th Ave. N.E. (northbound) to N.E. 150th St. Turn right onto N.E. 150th. After two blocks, the State Laboratories are on your left.

**Note:** All laboratory samples, specimens and supplies must be taken to the PHL loading dock near the center of the building. No deliveries will be accepted in the reception area at the main entrance.

## Driving Map



## Campus Map





## **COLLECTION AND SUBMISSION OF SAMPLES AND SPECIMENS**

### **SAMPLING & SPECIMEN COLLECTION KITS PROVIDED BY PHL**

#### **Specimen Kit Requisition Policy**

To order write:      Washington State Public Health Laboratories  
                                 1610 N.E. 150th Street  
                                 Shoreline, WA 98155

With the first order, you will receive an order sheet for subsequent use. As supplies and kits shelf life are limited, plan to order no more than a month's supply. Mail Services telephone number is (206) 418-5579 and fax number is (206) 418-5405 or e-mail to [phl.mailroom@doh.wa.gov](mailto:phl.mailroom@doh.wa.gov)

IATA and postal regulations require the use of double mailing containers for submission of cultures and certain other material. When requesting mailing containers, please specify the type of culture (enteric, TB, etc.) so you will receive the appropriate kit and laboratory form. Most of the specimen kits use double mailers. Always wrap the laboratory form around the inner cardboard mailer, to avoid contamination if the specimen leaks.

When submitting a bacterial or viral isolate by any means of transportation, the package must be packed in agreement with IATA and USDOT regulations for Infectious Substances. The State Laboratories do not supply the packaging, but materials are commercially available from many sources. See Appendix A for the Federal Regulations, which apply to shipping etiologic agents.

**Please fill out the laboratory form completely.** Telephone numbers have been given for areas of the laboratories. Whenever questions arise regarding specimens or any of the services provided by the State Laboratories, a phone call is welcomed and will often save time and effort. Please print clearly when filling out all paperwork.

Mailing Kits Available		
KITS	CONTENTS	REMARKS
<b>Enteric Pathogens For more information call: Call (206) 418-5456</b>		
Stool or rectal	Cary-Blair transport medium, sterile swab, Enteric Bacteriology form, and instructions, inside a double cardboard mailer.	Use for isolation of enteric pathogens from stools: Salmonella, Shigella, E. coli, Yersinia, Vibrio and Campylobacter. Use sterile applicator swab to collect specimen, insert swab into Cary-Blair transport medium, break off stick at the score line below lid of bottle, push cap on <u>tightly</u> , seal with pressure-sensitive labeling tape and mail immediately.
Urine (Typhoid specimens only)	Buffered Glycerol Saline transport medium (pink solution), Enteric Bacteriology form, in a double cardboard mailer.	Add amount of specimen equal to volume of transport solution
Enteric pathogen cultures for identification	Enteric Bacteriology form, double cardboard mailer.	Pure cultures only, use screw-cap tubes, <u>do not mail</u> Petri plates (use courier service). <u>Do not send in liquid media</u> . Campylobacter jejuni cultures should be sent on blood or chocolate agar slants in screw-capped tubes. Salmonella, Shigella and Vibrio cholera confirmation is required by the Washington State Board of Health regulations.
<b>Tuberculosis For more information call: Call (206) 418-5473</b>		
Cultures for identification or drug susceptibility testing	Mycobacteriology form, double cardboard mailer	M. tuberculosis confirmation required by state law
Biopsy Material or Swab	Sputum kit, Mycobacteriology form, double cardboard mailer.	Keep specimen moist using a small amount of sterile distilled water or sterile saline.
Sputum	Centrifuge tube, Mycobacteriology form, directions, double cardboard mailer, whirl bag, and absorbent paper.	Single early morning specimens taken on 3 consecutive days (2-3 teaspoons per specimen)
Stool	Centrifuge tube, Mycobacteriology form, and double cardboard mailer.	Specimens must be received within 24-hours of collection, notify before shipping
Urine	Centrifuge tube, Mycobacteriology form, and double cardboard mailer.	Three single clean-catch early morning specimens taken on consecutive days (30 ml per specimen)
<b>Parasitology - For more information call: (206) 418-5469</b>		
Pinworms	Two (2) tubes with vaspar swabs, Parasitology form, directions, double cardboard mailer.	Collect a morning specimen on two successive days, before the patient uses the bathroom
Sputum specimen	Sputum in screw-cap tube or bottle with equal quantity of 10% formalin, Parasitology form, double cardboard mailer.	For examination for Paragonimus eggs or Strongyloides larvae
Stools	Para-Pak ULTRA <sup>®</sup> ECOFIX <sup>™</sup> Parasitology form, instructions, plastic bag, mailer	The Para-Pak ULTRA ECOFIX kit contains a non-formalin based preservative, which is necessary for parasites preservation. Specimens should be placed into this kit as soon as possible after collection. Fresh stool specimens are not recommended because of transport time. Observe expiration date on kit.
Urine		For examination of Schistosoma haematobium eggs, add 5 ml of 10% formalin to urine after collection. Eggs are most likely to be present in last few drops of urine, especially if urine contains blood or pus. Recommend repeating test 3 consecutive days.
<b>Special Bacteriological Pathogens (Reference) - For more information call: (206) 418-5452</b>		
Bacteriology Culture	Reference Bacteriology form, double cardboard mailer.	Viable pure culture. Do not mail Petri plate; Use a courier service. A valid attempt to identify the organism is required. Send laboratory results obtained.



Mailing Kits Available		
KITS	CONTENTS	REMARKS
Special Pathogens Clinical Specimen		See Special Bacteriological Pathogens Section for clinical specimens which will be accepted by the State Laboratories
Legionella cultures and Direct FA	Reference Bacteriology Legionella Culture- DFA form, double cardboard mailer.	Use a Reference Bacteriology Legionella Culture- DFA form for culture or Direct FA
Brucellosis, Tularemia	Serology-Bacterial, Fungal, Parasitic form, double cardboard mailer.	Febrile agglutination. Acute and convalescent specimens required. Contact the unit before submitting specimens.
Legionnaires' Disease Serology	Serology-Bacterial, Fungal, Parasitic form, double cardboard mailer.	Acute and convalescent specimens preferred
<b>Special Respiratory Pathogens - For more information call: (206) 418-5492</b>		
Group A Beta Hemolytic Streptococcus (Strep kits) Clinical cultures	Silica gel, Pai slant, swab, Nose and Throat form, double cardboard mailer.	Reference cultures accepted
Diphtheria Clinical specimens, contact or case	Two Pai media slants, Nasopharyngeal and throat swabs, Nose and Throat form, double cardboard mailer.	Take throat and nasopharyngeal cultures. Notify the Special Respiratory Pathogens Unit.
Diphtheria Culture	Nose and Throat form, double cardboard mailer.	Confirmation of C. diphtheria required by state law
Pertussis	Slides and holder, nasopharyngeal swab, charcoal transport media, Nose and Throat form, directions, double cardboard mailer.	Do not heat-fix slides. Diagnosis of pertussis requires both culture and smear to be taken. Reference cultures accepted.
<b>Sexually-transmitted Diseases - For more information call: (206) 418-5492</b>		
Gonorrhea Clinical specimens	Transport media plates with plastic bag and CO <sub>2</sub> tablets, swab, Fluorescence Microscopy form, send per shipping regulations.	Incubate the plates at 35° for 48 hours before sending to the State Laboratories.
Reference culture for confirmation	Fluorescence Microscopy form, double cardboard mailer for tubes or cardboard mailer for plates.	Submit viable pure culture. Incubate culture 24 hours before mailing.
<b>Virology - For more information call: (206) 418-5458</b>		
Virus Isolation	Viral transport medium, swab, and Virus Examinations form, double cardboard mailer.	Call (206) 418-5458 prior to sending samples. Ship in special mailing containers with ice packs. No wet ice.
Herpes	Lab form, VTM and swabs	Specimens accepted only from Local Health Jurisdictions, Family Planning and Planned Parenthood organizations.
Respiratory Virus Isolation	Viral transport media, swab, and Virus Examinations form, 2-1.8 ml sterile PBS tubes, sterile pipette and collection container, double cardboard mailer.	Call prior to sending samples for virus isolation. Ship in special mailing containers with ice packs. No wet ice.
Rabies	Rabies Laboratory Report and Animal History form, directions, special bio-transport shipping container and bag, absorbent, ice packs	Submit animal heads. Ship with ice packs. Send through your local health jurisdiction. Notify Virology Unit before shipping.
Virus	Virus and Rickettsial Examinations form, double cardboard mailer, directions.	Acute and convalescent specimens preferred. Consult unit for single specimen.
Influenza	Laboratory form, VTM and swabs, return box.	Accepted from surveillance physicians, nursing homes, all Local Health Jurisdictions, and others pre-approved by Department of Health Communicable Disease Epidemiology.
Rabies	Laboratory form, secondary and specimen	All Local Health Jurisdictions or by pre-approval of

Mailing Kits Available		
KITS	CONTENTS	REMARKS
	containers, absorbent material, mailer, outer box and ice packs.	Dept. of Health Communicable Disease Epidemiology.
<b>Water Microbiology- For more information call: (206) 418-5491</b>		
Drinking Water Environmental Water Pool or Spa Water	Sterile plastic bottle with sodium thiosulfate, Water Bacteriological Analysis form, instructions, mailing container, and return address label. Only kits supplied by PHL or local health jurisdiction will be tested.	Specimen must be in the laboratory within 30 hours after collection. Samples are accepted Monday through Thursday 8 a.m. to 5 p.m. and Fridays 8 a.m. to 12 p.m. Water Bacteriology kits <u>MUST</u> be prepaid. The fee is \$20.
<b>Food Bacteriology- For more information call: (206) 418-5442</b>		
Food	No kit provided	Call local health jurisdiction and CD Epidemiology (206) 418-5500 before sending food or specimen.
Stool	Food Bacteriology form, sterile cup, sterile sticks. for transport	Stool must be fresh. <u>Do not</u> use transport media. Call local health jurisdiction and CD Epidemiology (206) 418-5500.
<b>Newborn Screening- For more information call: (206) 418-5410</b>		
Newborn	Collection forms, parent information pamphlet, filter paper card, instructions for collection, and a protective envelope for mailing	Follow instructions on back of card
Whole Blood Testing	Dried Blood Hemoglobin form	Collection instructions on form

## SPECIMEN COLLECTION

### Clinical Specimen Collection:

The collection of clinical specimens must follow established laboratory policies and procedures. These policies and procedures must be documented as required by Chapter 246-338 WAC, Medical Test Site Rules, State of Washington Department of Health, Office of Laboratory Quality Assurance. Refer to the table below for general specimen submission instructions. Turn to the submission guidelines of each laboratory unit i.e. serology to which you will be sending the specimen for specific detailed information

SPECIMEN SUBMISSION INFORMATION
<b>Blood, serum or cerebral spinal fluid (CSF)</b>
<ol style="list-style-type: none"><li>1. Do not freeze whole blood.</li><li>2. Submit specimens in a screw-cap tube sealed with waterproof adhesive tape wound in the direction that tightens the cap. <i>If the tube leaks during shipment, we reserve the right not to test the specimen.</i></li><li>3. Place tubes in individual plastic zip-lock type bags. <i>Do not put the laboratory form in direct contact with the specimen tube.</i> Use sufficient absorbent material to secure the contents and contain any leakage during shipment.</li><li>4. Wrap the laboratory form around the outside of the plastic zip-lock type bag and secure with rubber bands. Do not include more than four (4) laboratory forms with each inner can. Place the plastic bag and the laboratory form into a cardboard mailing container. Finally, place the cardboard mailing container into a plastic biohazard bag. Note that more than one cardboard container can be placed into the plastic biohazard bags for shipment.</li><li>5. Place a mailing address label onto the plastic biohazard bag before shipping specimen(s). On the outside of the biohazard bag, please specify the nature of the specimen or contents (Diagnostic, blood sample, etc).</li></ol>
<b>Blood Spots, Dried (Newborn Screening Cards)</b>
<ol style="list-style-type: none"><li>1. Use the Newborn Screening card kits supplied by the PHL.</li><li>2. <b>Air-dry</b> the specimen at <b>room temperature</b> until thoroughly dry (minimum 2 hours).</li><li>3. Place specimen into the special envelope provided.</li><li>4. Do not enclose in plastic. This will INVALIDATE the specimen.</li><li>5. Within <b>24 hours</b> of collection, submit to the Newborn Screening Laboratory at the address on the front of the envelope.</li></ol>
<b>Cultures:</b>
<ol style="list-style-type: none"><li>1. Submit cultures to the PHL in screw-capped tubes only. <b>DO NOT</b> submit cultures in Petri dishes. All screw cap culture tubes must be tightly sealed to the tube with waterproof adhesive tape wound in the direction, which tightens the cap.</li><li>2. Place all culture tubes into plastic zip-lock type bags, and then package using approved shipping containers. Do not send cultures in office stationery envelopes or other non-approved containers.</li></ol>

<b>SPECIMEN SUBMISSION INFORMATION</b>	
<p><b>Note:</b> Always follow IATA and USDOT regulations (see Appendix A).</p> <p>3. <b><i>DO NOT</i></b> send broth cultures unless it is absolutely necessary. Contact the appropriate PHL laboratory unit prior to sending if shipment is necessary.</p>	
<b>Virology Specimens</b> (for isolation, serology or direct antigen testing)	
<p>1. Specimens are preferably received in the laboratory within 24 hours of collection. After 24 hours, the viability of viruses will decrease, which may result in the inability to detect virus.</p> <p>2. All specimens must be shipped cold - NOT FROZEN.</p> <p>3. Notify the Virology Laboratory when specimens are to be shipped - especially specimens to be shipped on Friday or Saturday.</p>	
<b>Rabies Specimens</b>	
<p>1. Animals suspected of having rabies <i>MUST</i> be referred to the local health jurisdiction for handling and shipping.</p> <p>2. The local health jurisdiction will notify the PHL's Virology Unit by phone when shipping specimens for rabies testing.</p>	
<b>Specimens to be tested at the Centers for Disease Control and Prevention (CDC)</b>	
<p>1. All Specimens being shipped to CDC in Atlanta, GA, must be routed through the PHL.</p> <p>2. Turn-around times for results on these specimens will vary. Contact the individual PHL unit for specific information.</p> <p>3. A CDC DASH form must be enclosed with each specimen forwarded to the CDC. Please contact the PHL to request CDC DASH forms.</p>	

## Environmental Samples:

The collection of environmental samples must follow established laboratory/field policies and procedures. These policies and procedures must be documented. Refer to the material below for general sample submission instructions. Turn to the submission guidelines of each laboratory unit (i.e. Inorganic Chemistry) to which you will be sending the sample for specific detailed information.

Call us at (206) 418-5400 if you have questions about samples, interpretations, procedures, or any other aspect of Public Health Laboratories services. For public health emergencies after hours, call Communicable Disease Epidemiology at (206) 418-5500.

Kits are expensive and many have expiration dates. Return all unused and outdated specimen kits and mailing containers to the PHL for recycling. For information regarding mailing containers, biohazard bags or media, call Mail Services at (206) 418-5579 or fax (206) 418-5405 or e-mail to [phl.mailroom@doh.wa.gov](mailto:phl.mailroom@doh.wa.gov).

## Submission Procedure

1. Complete the appropriate laboratory form specific to each PHL laboratory unit. The form must include patient ID, submitter name, mailing address and submitter phone number, and date of collection.
  - a. Use *black, non-smearing ink* and please print clearly.
  - b. All specimen ID information must correspond with the laboratory form.
2. Include your name, return address, phone number, and date with all specimens, letters, memos and requests for laboratory supplies.
3. All specimens submitted to the PHL must have the return address of the submitter and the name of the person requesting the examination.
4. The PHL receives shipments from Greyhound, UPS, Federal Express, and the United States Postal Service Monday through Saturday. Call the PHL at (206) 418-5579 before sending samples/specimens.
5. If there is a laboratory fee required for testing, make check or money order payable to the Department of Health and send to the Department of Health, Revenue Section, and PO BOX 1099, OLYMPIA, WA 98507-1099. *Never send your payment with the specimen.*
6. Before sending specimens, make sure there is sufficient postage. The Postal Service will not deliver packages that do not have the required postage. *Do not send specimens collect* unless you have made prior arrangements with the PHL.

**Note:** Greyhound Express shipments are routinely picked up by PHL staff at about 6:30 a.m. each weekday. To accommodate other bus arrival times (such as emergency and other special arrangements), courier service will be provided for delivery to the PHL.

During regular business hours, please call the lab units involved to make special arrangements before sending the specimens. (See pages 4-7)

## HAND DELIVERY

Courier deliveries are received from 7:30 AM to 5:00 PM, Monday through Friday. The Public Health Laboratories are closed on weekends and holidays. Special arrangements must be made with laboratory personnel prior to delivery for any high priority items arriving outside the hours of normal operation.

All laboratory samples, specimens, and supplies must be taken to the PHL specimen receiving entrance, near the loading dock at the center of the building. No deliveries are accepted in the reception area at the main entrance. The loading dock is located past the main entrance in the middle of the building, indicated with signage. The glass door to the right of the loading dock has a doorbell for specimen delivery. Ring bell to summon mailroom staff to accept delivery. All delivery persons must have picture identification and will be required to sign the delivery log as shown below.

Specimen Sign-In Sheet							
Date	Time	Company / Courier	# of Pkgs	Sender	Specimen Type	Pick-up time	Employee Name

For questions, or to arrange delivery outside normal receiving hours, call the appropriate laboratory within PHL.

**Important:** When submitting specimens in person or by courier DO NOT leave the packages or specimens outside the building. All items must be delivered to an actual person.

## SHIPPING

**Important:**

*Appropriate regulations for the shipment of infectious materials must be followed when sending specimens to the PHL.* In conjunction with appropriate training, the following resources may be used for shipping and mailing regulations:

- [International Air Transport Association \(IATA\)](#) - Dangerous Goods Regulations 46<sup>th</sup> Ed. (1/1/2005-12/31/2005); <http://www.iata.org>; <http://www.iata.org/dangerousgoods/index>
- US Postal Service – Domestic Mail Manual Section C023; <http://pe.usps.gov/>– Title 39 Code of Federal Regulations Part 111
- US Dept of Transportation – Title 49 Code of Federal Regulations Parts 171-185; <http://www.myregs.com/dotrspa/>
- Appendix A of this directory contains more information on shipping requirements including shipping reference tables.

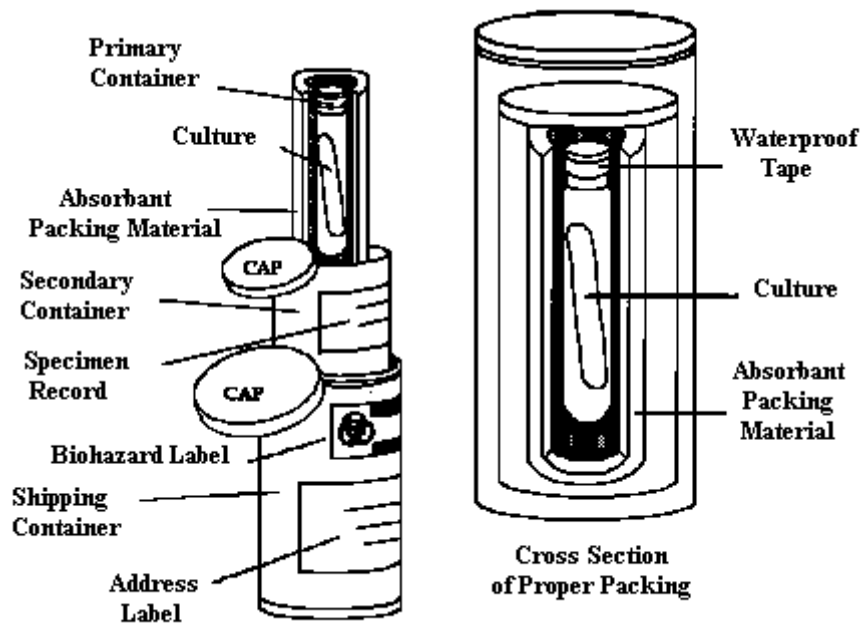
Shipping of diagnostic specimens and infectious materials should be performed or supervised by a person who has received training in the shipping of such materials. *It is the shipper's responsibility to ensure that packages being shipped meet current regulations.* The Code of Federal Regulations can be accessed at <http://www.gpoaccess.gov/cfr/index.html>. A copy of PHL shipping regulations may be obtained by contacting the PHL mailroom at (206) 418-5579. *Ice packs* must be used when submitting specimens in cooler boxes. In this directory *ice packs* refers to any one or a combination of: gel packs, frozen coolant packs, blue-ice packs, combination water and gel packs, or leak-proof plastic containers. It is important to ensure these products will not leak during shipment. Do NOT use wet ice to transport specimens to the PHL. *Leaking packages may be rejected.*

## INSTRUCTIONS FOR PACKING AND SHIPPING SPECIMENS

1. On the sample label print patient's complete name or confidential identifier corresponding with the requisition form, the type of specimen collected, and the date collected.
2. **Do Not** use ballpoint pens, wax, indelible pencils, or other writing instruments that tend to smear.
3. Enclose a completed laboratory request form with each and every properly labeled specimen, even if packaged in the same container.
4. Enclose the specimen in a screw cap tube or vial with a tight-fitting cap. For specimens shipped at ambient or higher temperatures positive means of ensuring a leak proof seal must be used, such as a skirted stopper, or metal crimp seal. If screw caps are used seal the cap with waterproof adhesive tape wound in the direction that tightens the cap.
5. Package specimens properly for transit (Figure 1) ensuring that personnel who handle the package will not come into contact with the enclosed specimen.
6. Place the tube or vial (primary container) in a watertight secondary container. Pack a suitable absorbent material around the tube to absorb shock and sufficient to absorb possible leakage of the entire contents. If several tubes are to be packed within the same can, wrap each tube individually in absorbent material. **Do not** place the request form within the secondary container; wrap it around the outside of the secondary container.
7. Place the secondary container into an outer shipping container. Seal the outer shipping container securely; affix a properly completed address label with a return address and postage, if required.
8. If specimens must be sent refrigerated or frozen, they should be packaged in an insulated container. The insulated container should be placed within a properly labeled cardboard box and sealed securely. The specimens should be packaged in a manner that prevents movement within the insulated container.
9. Try to time shipments (when possible) to arrive early in the week. Be particularly careful to avoid having the specimen arrive on a weekend or a holiday when possible. Call the PHL 24-hour number if shipments will be received outside of normal business hours. The 24-hour numbers are (206) 418-5500 and (877) 539-4344.
10. **Never** mail any clinical specimens or cultures in petri dishes.
11. Improperly packaged specimens and specimens that have leaked **may not be accepted.**
12. A specimen arriving with an incomplete or no request form may be held until the information is received. The proper request form for each specimen submitted must be

completed as fully as possible. When possible, include patient name or confidential ID, date of specimen collection, type of specimen, birth date or age, sex, date of onset, diagnosis, symptoms, attending physician, county of residence, suspected agent, reference culture information including type of medium and source of isolate, and other pertinent medical information including contact with insects, animals, etc., antibiotic or anti-tuberculosis therapies, recent vaccinations, similar infections in the family or community, and recent travel including destination and dates.

13. Copies of the reports are mailed only to the source indicated on the request form. Be sure to include the full 9-digit zip code for each address.



**FIGURE 1\***

\* 42 CFR, Part 72  
CDC Instructions  
Reference & Disease Surveillance  
Center for Infectious Diseases, Feb. 1986  
Office of Biosafety, Centers for Disease Control  
Atlanta GA 30333



## CONFIDENTIALITY

The Public Health Laboratories (PHL) places a very strong emphasis on protection of confidential data. The PHL also places a similar emphasis on providing timely results. In an attempt to ensure that these goals are met, the PHL requests that providers sign and return a Fax Confidentiality Statement stating that the receiving fax machine at the provider's facility is in a secure location and that only authorized personnel have access to faxed information. A sample of the Confidentiality Notice that will accompany each fax is provided below.



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**Public Health Laboratories**  
1610 NE 150<sup>th</sup> Street  
Shoreline, Washington 98155-7224  
Phone (206) 418-5400 FAX (206) 418-5445

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To: \_\_\_\_\_

Date: \_\_\_\_\_ Time: \_\_\_\_\_

Fax No. (\_\_\_\_\_)\_\_\_\_\_-\_\_\_\_\_

From: (Name)\_\_\_\_\_

Phone No: (\_\_\_\_\_)\_\_\_\_\_-\_\_\_\_\_

Number of pages: \_\_\_\_\_+ cover sheet

Message:

## **OFFICE OF ENVIRONMENTAL LABORATORY SCIENCES**

This Office provides testing services for microbiological, chemical, and radiological analyses of drinking water and a wide variety of environmental sample types, including food and shellfish in order to determine any potentially harmful health effects from the environment or environmental contamination.

This unit, in support of the Department of Health programs, performs the majority of the chemical analyses. The Department of Ecology, the Department of Agriculture, local health jurisdictions and private citizens utilize these laboratory services as well.

The laboratories of this Office serve as the reference laboratories for bacteriological and radiological tests in drinking water, food pathogen tests, and parasitology tests in support of State Environmental Health Programs.

To confirm the ability of the laboratories to perform analyses and serve as a reference facility, the laboratories staff participate in proficiency testing sponsored by the Federal Food and Drug Administration (FDA), College of American Pathologists (CAP), Center for Disease Control and Prevention (CDC), Federal Department of Energy's Quality Assessment Programs (QAP), Department of Ecology (DOE) Mixed Analyses Performance Evaluation Program (MAPEP), and activities of the Northwest Regional Quality Assurance Task Force. The laboratories also use private laboratories certified by NIST as proficiency test providers.

## **CHEMICAL TERRORISM RESPONSE**

The Chemical Terrorism Response Unit provides testing services of human blood and urine specimens for 20 heavy metals, cyanide, and volatile organic chemicals (VOCs) in support of the Washington State Emergency Preparedness and Response Program. Services are provided ONLY to local, state and federal health jurisdictions and law enforcement. State-of-the-art instrumentation and methods are used to identify and quantify exposure levels. The testing methodology varies according to the origin of the specimen and the type of chemical agent suspected or known to be involved.

Specimen collection information is available by contacting (206) 418-5643 or (360) 236-3387.

CHEMICAL AGENTS SURVEILLANCE				
Collection and Submission Instructions				
AGENT	SPECIMEN	COLLECTION TIME	TRANSPORT TEMP.	REMARKS
Suspected chemical agent exposure	Blood	Call 206-418-5643 or 206-418-5476	Refrigerate at 4°C	Prior approval by CT Response Unit required: 360-236-3387 or 206-418-5476. Ship promptly. Notify CT Response Unit at when and how specimens are being shipped.
Suspected chemical agent exposure	Urine	Call 206-418-5643 or 206-418-5476	Flash freeze at -70°C or on dry ice and keep frozen at -20°C or colder	Prior approval by CT Response Unit required 360-236-3387 or 206-418-5476. Ship promptly. Notify CT Response Unit at when and how specimens are being shipped.

#### Turn-Around Times

No turn-around times are available.

## FOOD MICROBIOLOGY

The Food Microbiology Unit examines specimens from suspect food-borne illness episodes to determine sanitary quality and to isolate and identify possible etiological agent(s). *It is **very important** that the CD Epidemiology Section and the Food Unit be notified prior to specimens being shipped.*

#### Turn-Around Times (working days):

Food..... 7 - 10 days\*  
 Environmental..... 7 - 10 days  
 Stools & Vomitus\*.....7 - 10 days\*

\* Listeria testing takes up to 30 days to complete.

#### Food testing guidelines

The Washington State Department of Health (WDOH) Public Health Laboratories (PHL) provide diagnostic services that include food microbiology. As a general rule, two categories of food testing are performed at the PHL:

---

\*\* Vomit must be neutralized within 1 hour of discharge (release) (episode)

### ***A. Food safety testing***

The Food Microbiology Laboratory conducts food safety testing at the request of local health jurisdictions based on inspections or complaints. Tests to determine water activity, pH levels, standard plate counts, coliform counts, and fecal coliform counts are performed. Approval from Food Microbiology staff (206-418-5469) must be obtained before specimens are submitted for food safety testing.

### ***B. Testing for etiological agents of public health and epidemiological concern***

In general, food tests for etiological agents of public health and epidemiological concern at the PHL are conducted *in outbreak\* situations only, and only after human specimens have been analyzed with positive results*. If, in an outbreak situation, there is leftover food, it is always advisable to collect food samples and hold them in case laboratory testing of food by the PHL is warranted. **However, food should not be submitted to the PHL without the approval of Communicable Disease Epidemiology (CDE).** Call 206-418-5500 (24 hour telephone number) if you would like to discuss food testing with CDE staff. Any exception to the outbreak/confirmed clinical results criteria must be approved by CDE staff. Food will not be accepted or tested for etiological agents without CDE approval.

*(\*A foodborne outbreak is defined as an incident in which: (a) two or more persons experience a similar illness, usually gastrointestinal, after ingestion of a common food, AND (b) epidemiologic evidence and/or laboratory testing evidence implicate a common food as the source of the illness. A single case of botulism always warrants food testing.)*

Samples must be clearly and completely identified. Include a WDOH PHL Food Specimen form with each sample. Forms can be obtained by calling the PHL Mailroom at 206-418-5579. The following information is considered necessary:

- Sample description
- Collector's name
- Name and address of the manufacturer
- Lot number
- Dealer or distributor
- Date, place, time of collection
- The reason for testing

In general, leave the food sample in the original container if available. Give each sample a securely attached label. Keep frozen, if frozen. Keep cold if the sample is already refrigerated or if the sample is at risk of spoiling during shipment, otherwise ship at ambient temperature.

**GUIDELINES FOR SUBMITTING FOOD AND FOOD-RELATED SPECIMENS TO  
THE WDOH PHL**

<b>AGENT/ DISEASE</b>	<b>SPECIMEN &amp; QUANTITY</b>	<b>COLLECTION TIME</b>	<b>TRANSPORT CONTAINER</b>	<b>REMARKS</b>
<i>Bacillus cereus</i>	Stool – Walnut-size (50 gm).	At onset.	Clean container or cup, keep cold, no transport media.	Isolation from stool alone does not confirm foodborne illness. Meaningful only if there is isolation from a food sample also.
	Food – 100 gm (1/4 lb).	At onset.	Clean container, keep cold.	
<i>Clostridium perfringens</i>	Stool – Walnut-size (50 gm).	At onset.	Clean cup or container, refrigerate, keep cold, no transport media.	Isolation from stool alone does not confirm foodborne illness. Meaningful only if there is isolation from a food sample also.
	Food – 100 gm (1/4 lb).	At onset.	Clean container, keep cold (10°C). Temperature is critical, rapid die-off of vegetative cells occurs below 10°C, spores are unaffected.	
<i>Clostridium botulinum</i>	Food – food remnants, washed/unwashed food container.	At onset.	Do not freeze. Ship food and food remnants in clean, leak-proof container; place in plastic bag (with absorbent material), then in an insulated shipping container. Ship cold with ice pack.	Notify Special Bacteriological Pathogens Unit (ph: 206-418-5452).
	Serum – 10 to 15 ml.	Collected soon after onset of symptoms and before antitoxin is given.	Place specimen in a clean, leak proof container; place in plastic bag, then in an insulated shipping container. Ship cold.	For all <i>Clostridium botulinum</i> specimens prior approval by WDOH Communicable Disease Epidemiology is required. Call 206-418-5500, 24 hours per day.
	Vomit – 10 to 15 ml.	At onset.	Ship same as serum.	For most <i>Clostridium botulinum</i> specimens, unless otherwise stated, notify Special Bacteriological Pathogens Unit as to when and how specimens are being shipped. Call 206-418-5452.
	Gastric material – Walnut-size (50 gm).	At onset.	Ship same as serum.	
	Stool – 10 to 50 gm. Enema material is acceptable. Obtain specimen from sterile (non-bacteriostatic) water enema. A volume of 20 ml collected after enema is sufficient.	At onset.	Ship same as serum.	
<i>Escherichia coli</i> O157:H7, other enterohemorrhagic strains of <i>E. coli</i>	Stool – Swab coated with stool specimen or rectal swab.	At onset.	Cary-Blair transport media, sent in double mailer.	Send isolate to the PHL Enteric laboratory.
	Food – 100 gm (1/4 lb).	At onset.	Sterile container, keep cold.	
<i>Salmonella</i>	Stool – Swab coated with stool specimen or rectal swab.	At onset.	Cary-Blair transport media, sent in double mailer.	Send to Enteric Lab (206-418-5456).
	Food – 100 gm (1/4 lb).	At onset.	Clean container, keep cold.	

AGENT/ DISEASE	SPECIMEN & QUANTITY	COLLECTION TIME	TRANSPORT CONTAINER	REMARKS
	Drinking water – one gallon.	During outbreak.	Clean jar, keep cold.	
	Environmental swabs.	At onset.	Ambient temperature.	Contact and transport directly to Food Microbiology unit.
<b>Shigella</b>	Stool – Swab coated with stool specimen or rectal swab.	At onset.	Cary-Blair transport media, sent in double mailer.	Send to Enteric Lab (206-418-5456).
	Food – 100 gm (1/4 lb).	At onset.	Clean container, keep cold.	Organism is difficult to isolate from food.
<i>Staphylococcus aureus</i>	Stool – Walnut-size (50 gm).	At onset.	Clean container, keep cold.	Isolation from stool alone does not confirm foodborne illness. Meaningful only if there is isolation from a food sample also.
	Food – 100 gm (1/4 lb).	At onset.	Clean container, keep cold.	Interpretation of the results depends on type of food and food handling.
	Enterotoxin studies/Food – minimum 100 gm (+100 gm above).	At onset.	Clean container, keep frozen.	Send to FDA
	Vomitus – Check pH when collected. Neutralize with sodium hydroxide or baking soda, if necessary.	During symptoms	Clean container, keep cold.	Must neutralize (pH) immediately after collection and must be received in Food Microbiology Unit within a few hours after collection.
Parasites	Stool – Walnut size.	At onset.	Use Ova & Parasite kit. Contact PHL Mailroom for kit (206-418-5579).	Consult with Communicable Disease Epidemiology (206-418-5500).
	Food – Currently (2/2006) food is not tested for parasites at the PHL.			Consult with Food Microbiologist (206-418-5442).
<i>Vibrio parahaemolyticus</i>	Stool – Swab coated with stool specimen or rectal swab.	At onset.	Cary-Blair transport media, sent in double mailer.	Specify test on Enteric form.
	Food – 100 gm (1/4 lb).	At onset.	Clean container, keep cold.	Do not put food directly on ice pack. Insulate with newspaper.
Viral (Norwalk-like)	Food – Currently (2/2006) food is not tested for parasites at the PHL.			
Chemical	No food or stool testing.			Consult with CD Epidemiology (206-418-5500).
Other Tests: Water Activity PH	Food – 100 gm (1/4 lb).	No priority.	Clean container.	Ambient temperature, or if cold, keep cold.
	Food – 100 gm (1/4 lb).	At onset.	Clean container.	Ambient temperature, of if cold, keep cold.

1. Food samples should be kept cold, but not frozen, during shipment. A styrofoam cooler with ice packs is recommended for protecting the sample during the shipping process.

2. Food samples must be protected from direct contact with the ice. Placing insulation between the sample and the ice pack will provide protection. Newspaper, bubble wrap, or other padding material can be used for the insulation.
3. Samples should be transported to the laboratory as soon as possible for best results. Samples may be shipped via Greyhound bus, Fed Ex, UPS Express or hand delivered immediately after collection.
4. Food samples are accepted Monday through Thursday from Friday 7:00am to 5:00pm. Special deliveries on Saturday can be prearranged by contacting the food laboratory at (206) 418-5442.

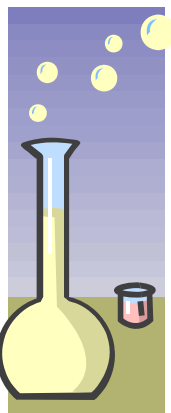


## INORGANIC CHEMISTRY

Tests in this unit fulfill the requirements of the Environmental Protection Agency (EPA) and Washington State Drinking Water Standards for Inorganic Compounds (IOC's). Types of tests offered include:

REGULATED INORGANIC CHEMICALS		
Primary Chemicals (EPA)		
Antimony	Chromium	Nitrate
Arsenic	Mercury	Nitrite
Barium	Nickel	Selenium
Beryllium	Cyanide	Sodium
Cadmium	Fluoride	Thallium
Secondary Chemicals (EPA)		
Chloride	Manganese	Sulfate
Iron	Silver	Zinc
Complete Inorganic Test Chemicals (in addition to all listed above) (WA)		
Color	Hardness	Turbidity
Conductivity	Total Dissolved Solids	
Lead And Copper Rule (EPA)		
Copper		
Lead		

The "complete inorganic chemistry analysis" includes the parameters required by the EPA and the State of Washington. Detailed EPA and Washington State requirements are available from the Department of Health, Office of Drinking Water, within the Division of Environmental Health Programs, at (360) 236-3100.



The Inorganic Chemistry Unit can analyze for each of these items individually or as part of a total test package. EPA regulations require that all items marked in the Primary Chemicals list above be tested when meeting the requirements of a complete inorganic test. Washington State regulations may require the tests indicated in order to meet the requirement of a complete inorganic test.

The unit also tests for aluminum, cobalt, molybdenum, strontium, titanium, vanadium, and for the parameters of alkalinity, silica, residual chlorine, surfactants (MBAS), phosphates, ammonia, bromate, chlorate, chlorite, and bromide in water as well as lead in blood.

Sample container and information kits are available by calling (206) 418-5492.

This laboratory also performs lead tests in blood.

## Turn-Around Times

IOC – complete .....4-6 weeks  
 Metals.....4 weeks  
 Incomplete – primary.....4 weeks  
 Nitrates .....2 weeks  
 Fluorides.....2-3 weeks  
 Lead in Blood.....1-5 days

<b>KITS AVAILABLE DIRECTLY FROM THE TESTING LABORATORIES</b>		
<b>KITS</b>	<b>CONTENTS</b>	<b>REMARKS</b>
<b>Inorganic Chemistry- For more information call: (206) 418-5492 for fee/kit information</b>		
Complete Inorganic Chemicals in Water	Two 1-quart containers, mailing container, form, ice pack	As required for compliance with Federal and state regulations. Special parameters or groups of tests available. Must use container provided.
Fluorides in Water	125 ml bottle, mailing container, form	
Nitrate in Water	60 ml (2 oz) ice pack, form, bottle, mailing container	
Lead in Water		Call for fee and/or kit information

INORGANIC CHEMISTRY (IOC)						
Collection and Submission Instructions						
TEST	TRANSPORT CONTAINER	STORAGE TEMPERATURE	ANALYSIS TIME	SAMPLING & MAILING DAYS	SAMPLE LOCATION	RESULTS
Complete IOC Drinking Water	Two 1-quart cubitainers per sampling point/source, ice pack, styrofoam mailing container	Refrigeration at 4°C	48 hours for NO <sub>2</sub> and NO <sub>3</sub> tests	Monday - Wednesday Mail within 8 hrs of collection. Sample must be in lab within 24 hrs & received by Thursday.	Closest to source before tanks and/or treatment	Monday - Friday
Incomplete IOC Primary	Same as above	Same as above	Same as above	Same as above	Same as above	Monday - Friday
Metals	One 1-qt acid-rinsed cubitainer	Acidify (1% HNO <sub>3</sub> upon receipt)	6 weeks	Monday – Friday	Same as above, but if testing for lead or other metals leaching from plumbing, take sample 1st thing in AM, from COLD tap. Do not flush the system.	Monday - Friday
Nitrates / Nitrites	One 2-oz (60 ml) polypropylene bottle, ice pack, styrofoam mailing container	Refrigerate at 4°C	2 weeks	Monday – Wednesday	Closest to source; before tank &/or treatment	Thursdays
Fluorides	One 4-oz (125 ml) polypropylene bottle	Ambient	48 hours for non-chlorinated water 14 days for chlorinated water	Monday – Friday	Same as above - For fluoridated systems, AFTER fluoridation	Fridays
Lead and other trace metals in Blood	royal blue top (trace metal) Vacutainers	Refrigerate upon receipt		<b>CALL AHEAD -</b> Performed on a restricted basis. Submitter must supply container.  See Specimen Collection Section of this manual for blood.		
Others - Without Metals	One 1-qt cubitainer	Check with Lab for details	Check with Lab for details	Monday - Wednesday	Closest to source; before tank &/or treatment	Monday - Friday

## MARINE BIOTOXINS

The Marine Biotoxin Laboratory provides testing services of shellfish for Paralytic Shellfish Poisoning (PSP) and Domoic Acid, sometimes referred to as Amnesic Poisoning (ASP) toxins, in support of the State of Washington Food and Shellfish Program. The Office of this Program in Olympia (360) 4245 arranges collection of samples for PSP or Domoic Acid. Questions regarding sample collection and shipment should be directed to this office as well.



Shellfish  
Safety  
236-  
analysis.  
directed

### Shellfish Related Illness

Questions concerning illness associated with eating shellfish should be directed either to the local health jurisdiction or to the Office of Shellfish Programs (360) 236-3330 or to Communicable Disease Epidemiology (206) 418-5500.

### Sample Collection

1. Samples must yield at least 120 grams of meat for processing (approx. 4 oz.). Shellfish must be in the shell and fresh.
2. Do not submit specimens with cracked or crushed shells.
3. Place shellfish in a waterproof plastic bag.
4. Do not mix species (only one species per bag).
5. Sample size is according to the following table (assuming medium size):

MARINE BIOTOXINS	
Shellfish Collection Guide	
Species	Number* to Submit
Oysters	8 - 10
Butter Clams	8 - 10
Littleneck, Manila clams	30 - 40
Horse Clams	3 - 4
Eastern Softshells	10 - 12
Blue Mussels	75 -100
California Mussels	5 - 6
Rock Scallops	2 - 3
Pink Scallops	30 - 40
Geoduck	3 - 6
*These numbers are based on average-size specimens.	

### Handling

1. Do not hold shellfish in seawater or freshwater at any time after collection, however, rinsing with seawater or freshwater to remove sediment is recommended.
2. If samples must be held prior to shipment, keep refrigerated. Putrid or decomposed samples will not be processed.

### **Data Forms**

1. Complete the form using black ink.
2. Fill in dates collected and submitted, species and number of organisms.

### **Shipment**

1. Place samples in a container that can be sealed, such as a cardboard box. Plastic buckets are available upon request from the Office of Shellfish Programs at (360) 236-3330.
2. Frozen ice packs should be used to refrigerate samples during shipment.
3. Include one Shellfish Biotoxin Sample Form for each sample submitted. Place form in a separate waterproof plastic bag.
4. Submit samples as early in the week as possible. Samples can be shipped UPS or Greyhound. Call the Office of Shellfish Programs at (360) 236-3330 for other transportation arrangements.

### **Turn-Around Times**

PSP Results .....24 hours  
Domoic Acid Results.....2 days



## PARASITOLOGY

The Parasitology Laboratory provides clinical laboratory services **ONLY** to local health jurisdictions. This unit, however, serves as a reference laboratory for all laboratories within the state.

Reference services are available from the Centers for Disease Control and Prevention (CDC) for serology and identification of unusual parasites. If a parasitic disease is suspected from a symptomatic patient who has traveled in endemic areas, please contact the Parasitology Unit for specific instructions.

### Turn-Around Times

Reference, Gross, Ova Parasites Specimens ..... 2 - 4 days

**Note:** In events where further identification is needed, Turn-Around Times will vary.

PARASITOLOGY						
Collection and Submission Instructions						
AGENT	SPECIMEN	COLLECTION TIME	TRANSPORT CONTAINER	TRANSPORT TEMP	STORAGE TEMP	REMARKS
Arthropods	Suspected of causing human illness		Bottle containing tap water or 0.85% saline	Ambient	Ambient	*Ship in double cardboard mailer
Blood Parasites	Thick & thin blood smear stained	During fever and chills	Slide mailer and protective packaging (bubble wrap or peanuts)	Ambient	Ambient	Fix and stain slides immediately
Gross Parasites	Suspected parasite		Bottle containing water or 0.85% saline	Ambient	Ambient	Gross parasites from human patients only, wash adult worms free of stool, ship*.
Pinworms	Anal swab	Two successive mornings	Two Vaspar coated swabs in tubes	Ambient	Ambient	Brush swab lightly over anal area, then insert ¼ inch into anal canal, place swab in tube, ship*.
Protozoan cysts and Trophozoite  Helminth ova	Fresh walnut-sized stool (50 gm)	Every other day for three collections or three over a 10-day period	Para Pac ULTRA ECOFIX or one tube with 10% formalin, one tube with PVA	Ambient	Ambient	Make sure the patient adds enough stool specimen to the collection kit to reach the red line marked on the outside of the tube. Mix well. Use PHL collection kit for specimen and ship*.

# RADIATION

## Drinking Water Sample Analysis

The Radiation Unit is certified by the Environmental Protection Agency for testing of Drinking Water using the following methods:

	<u>EPA Method Code</u>
Gross Alpha & Gross Beta	900
Photon Emitters (including Iodine 131)	901
Radium-226	903
Strontium 89 & Strontium 90	905
Tritium	906
Natural Uranium	908
Radium-228	In-house procedure



The unit also performs radon analysis for drinking water.

## All Other Sample Types

The unit is capable of providing radiochemical analysis of almost any sample type. The Laboratory routinely tests soils, sediments, shellfish, fish, meat, sludge, mill tailings, milk, water, air, vegetation, and food products.

## Turn-Around Times

Wipes.....Per customer request. Can be as quick as 24 hrs.

Gross Alpha/Beta in Air .....	3 weeks
Gross Alpha/Beta in Water .....	4 weeks
Radium in Water .....	6 weeks
Radon .....	1 week
Uranium in Water.....	2 weeks
Uranium in Soil .....	4 weeks
Gamma in Milk, Water, Food, or Air.....	2 weeks
Gamma in Soil.....	4 weeks
Strontium in Water or Milk.....	5 weeks
Strontium in Air, Food, or Soil .....	8 weeks
Plutonium in Water .....	3 weeks
Plutonium in Soil, Food, or .....	4 weeks

The unit can meet quicker response times if necessary. Coordination of an increased priority will be made with all the programs that the laboratory supports. This is necessary, since for one set of samples to have a priority, another set of samples will likely experience an increase in turn-around times.

## Collection and Submission Instructions

The unit provides sample container and sampling information kits. For kit inquiries, call (206) 418-5486. The submitter will need to furnish all the information requested on the laboratory forms that are provided with the kits.

<b>Radiation Chemistry - For more information call: (206) 418-5498</b>		
Water	Container depends on type of chemical tests requested	Contact the Radiation Chemistry Unit at (206) 418-5484. Testing as required by the Federal Safe Drinking Water Act.
Environmental Samples [Other than Safe Drinking Water Act (SDWA) samples]		Contact the number listed above for detailed information

## **WATER MICROBIOLOGY**

The Water Microbiology Laboratory serves as a testing, reference and training resource for the State of Washington's Environmental Health Drinking Water and Food Safety and Shellfish Programs. The unit performs EPA and FDA approved methods in the interest of maintaining the safety of public drinking water supplies and environmental waters throughout the state.



As the Primary Drinking Water Microbiology Laboratory for the State of Washington, a monthly rotation schedule has been implemented to provide expertise in all approved EPA methods to meet primacy laboratory requirements. If a specific method is needed, please consult Water Microbiology personnel prior to sample submission, (206) 418-5491. Large-scale studies, other than routine sampling, should be arranged in advance with the Water Microbiology Laboratory.

## **DRINKING WATER MICROBIOLOGY**

The types of samples accepted include: routine drinking water, raw source water, swimming pools, spas, and other recreational waters. The following is a list of Drinking Water Microbiology tests available:



<b>METHODOLOGY LIST BY REGULATION</b>	
<b>Total Coliform Rule</b>  (Total Coliforms and Fecal Coliforms or E.coli)	<b>Surface Water Treatment Rule</b>  (Total Coliforms, Fecal Coliforms, and Heterotrophic Bacteria)
Chromogenic/Fluorogenic Methods  (Colilert and Colisure)	Membrane Filter (MF)
Membrane Filter (MF)	Multiple Tube Fermentation (MTF)
Multiple Tube Fermentation (MTF)	Heterotrophic Plate Count (HPC)
Single Volume Fermentation (PA)	

### **Drinking Water Bacteriology Kits**

Drinking Water Microbiology Kits are available for a fee of \$20.00 each. The fee charged includes both the kit and the analysis and must be purchased in advance of sample collection. The drinking water sample collection kit includes a special sampling bottle, laboratory form with instructions, and mailing container. Kits can be purchased in person or by mail using a check or money order. Please contact the billing department regarding the purchase of sample collection kits at (206) 418-5400.

### **Sample Requirements**

1. Routine drinking water sample collection must be less than 30 hours old when received in the laboratory for testing. Samples over 30 hours old become unsuitable and will not be tested.
2. Samples for routine testing are accepted Monday through Thursday from 7:00am to 4:00pm, and on Friday from 7:00am until noon.
3. Raw source waters, pool or spas and special requests for HPC testing must be less than 8 hours old when received in the laboratory for testing. Samples received after 8 hours will be tested but may produce invalid results. HPC requests are accepted Monday through Thursday from 7:00am to 12 noon.

## **Submission of Drinking Water Samples**

1. Use only the special sample collection kits furnished by the Public Health Laboratories.
2. The sample information form included in a sample collection kit must be filled out completely in accordance with the instructions printed on the back of the form.
3. It is recommended that Drinking Water samples be kept cold, but not frozen, during shipment. Using of blue ice packs with some insulation between the sample and the ice pack (i.e.: newspaper, bubble wrap, etc.) is recommended.
4. Samples must be less than 30 hours old when received in the laboratory for testing. Note: For raw source water or HPC test requests, samples must be received in the water lab within 8 hours of collection.
5. Samples may be mailed (Fed Ex, UPS Express) or hand delivered immediately after collection.
6. Routine drinking water samples are accepted Monday through Thursday from 7:00am to 4:00pm and on Friday from 7:00am until 12 noon (within 30 hours of collection).
7. Raw Source Waters, Pools or Spas, and HPC requests are accepted Monday through Thursday from 7:00am to 12 noon (within 8 hours of collection).

### **Turn-Around Times**

Results ..... 1 – 7 business days

## **MARINE WATER MICROBIOLOGY**

Marine Water testing requests related to the monitoring of shellfish growing areas must be coordinated through the Food Safety and Shellfish Program, (360) 236-3319. Process Water testing related to the exportation of shellfish goods is also available for a fee of \$21.00 per test. Prior arrangements must be coordinated through the Food Safety and Shellfish Inspection Program Lead, (360) 236-3313.

The following is a list of Marine Water Microbiology testing available:

- Growing Area Survey and Classification
- Fecal Coliform (MTF)
- Total Coliform and Fecal Coliforms (MTF)
- Wet Storage

## **Submission of Marine Water or Process Water Samples**

1. All samples must be co-coordinated through the Food Safety and Shellfish Program prior to submission.
2. Use only those water bottles furnished by the Water Microbiology Laboratory or the Food Safety and Shellfish Program.
3. Survey form(s) must be filled out completely and submitted with samples.
4. All Marine Water or Process Water samples must be shipped cold, but not frozen.
5. A temperature control bottle labeled "TC" must be included with each set of sample in order to verify that the samples remain between 0-10°C during shipment. Samples with temperature control bottles greater than 10°C are unsuitable and will not be tested.
6. Samples must be less than 30 hours old when received in the laboratory.

### **Turn-Around Times**

Results.....1 – 7 business days

### **Process Water Billing**

Process Water Billing invoices are prepared monthly. Please contact the Accounting Department with inquiries regarding Process Water testing.

Accounting Department  
Public Health Laboratories  
1610 N.E. 150th Street  
Shoreline, Washington 98155-9701  
(206) 418-5424

## **OTHER WATER MICROBIOLOGY TESTING**

### **Sterilization Monitor Test**

The Water Microbiology Laboratory processes Sterilization Monitor Tests to determine the effectiveness of Steam, Dry Heat, and Gas (Ethylene Oxide) sterilization. The submitter must obtain biological sterility indicators. A fee of \$10.00 is charged for each test set, two tests and one positive control. Contact the Water Microbiology personnel with inquiries regarding Sterilization Monitor Tests (206) 418-5491.

### **Giardia and Cryptosporidium**

The Water Microbiology Laboratory may provide Giardia and Cryptosporidium testing in response to outbreak situations. Please contact Water Microbiology personnel (206) 418-5491.

## WATER MICROBIOLOGY

### Sample Collection and Submission Instructions

Tests	Samples	Collection	Transport Container	Storage Temperature	Remarks
Drinking Water Samples For Total Coliforms and Fecal Coliforms or E.coli	120 ml of Drinking Water, Raw Source Water, Pools or Spas	Location of interest (sites which are representative of water quality throughout the distribution system)	Submit sample bottles in styrofoam container with ice or ice packs. Sample information forms should be packaged in a watertight ziploc bag along with samples.	Cold shipment of 0-10 °C is recommended. Do not store. Ship immediately. Drinking Water must be less than 30 hours old when received. Raw Source Water and Pools or Spa sample must be received in less than 8 hours.	Coliform tests use MTF, MF, C/F, MMO-MUG and P/A techniques appropriate to the type of sample. Include any special instructions with sample. Fecal coliform tests will be provided on any sample tested for total coliforms.
Heterotrophic Plate Count	120 ml of Drinking Water, Raw Source Water, and Pools or Spas	Location of interest	Submit sample bottle and form in a cardboard mailing container inside a biohazard bag.	Do not store. Ship immediately. Sample must be received in less than 8 hours.	Provided for all swimming areas. Also provided for other samples upon request.
Marine Water and Process Waters For Total Coliforms Fecal Coliforms or E.coli	120 ml of Marine Water or Process Water	Location of interest	Submit sample bottles in styrofoam container with ice or ice packs. Survey forms should be packaged in a watertight zip-loc bag along with samples.	Sample must be cold but not frozen. Maintain 0- 0-10°C. Do not store. Ship immediately. Marine Water and Process Waters must be less than 30 hours old when received.	Prevent samples from becoming submerged in melted ice. Submit a Temperature Control blank marked "TC" with each package.
Biological Sterility Indicators	Spore Strip Or Ampoule	Autoclave, Dry Heat, or Gas (Ethylene Oxide) sterilization	If liquid, wrap in absorbent material. Place tests in plastic watertight bag, ship samples and form in accordance with shipping regulations.	Ambient	Process in conjunction with a normal sterilizing run. Follow manufacturer's instructions. A \$10.00 fee is charged for testing. The submitter must obtain Biological Sterility Indicators.

### Special Tests

Special tests include: Microscopic Examinations, *Pseudomonas*, *Giardia* and *Cryptosporidium*. Consult with the Water Microbiology Laboratory for specific information regarding special testing.

# OFFICE OF LABORATORY OPERATIONS AND TECHNICAL SUPPORT

This office provides internal technical and operational support to the State Public Health Laboratories. Included within the office are, Technology Transfer, Media and Glassware Preparation, Mail Services, Fiscal Management, Instrument Maintenance and Facilities Maintenance.

## Central Services

Mailroom  
Media Preparation  
Glassware Preparation  
Specimen kit preparation  
Shipping and receiving

## Technology Transfer

Laboratory Training  
Meetings & Conferences

## Maintenance

Building and Grounds  
Motor Pool  
Security

Consultation from these areas is offered to local public and private health facilities. Areas of expertise include laboratory training, maintenance of laboratory equipment, facilities management, specimen handling, preparation of culture media, and shipping regulations

This office provides all the kits and containers used to deliver specimens to the State Laboratories and are responsible for their contents, quality control and shipping. During outbreaks of disease, laboratory support from this unit is coordinated with the efforts of local health officers, physicians, and state epidemiologists.

## **PHL Mailroom**

The PHL mailroom receives all mail, samples and specimens that are sent to the PHL. This unit also is responsible for preparing and supplying kits for many of the tests performed at the PHL. See section on collection and submission for details on submitting samples or specimens to the PHL.

## **PHL Maintenance**

The maintenance department is responsible for the upkeep of the PHL building and grounds, care of the cars in the motor pool, oversight on preventative maintenance of laboratory equipment, meeting room setup, building security and of the janitorial services.

## **Glassware and Media Preparation**

This department makes almost all of the media used by the PHL testing units. They are responsible for laboratory glassware preparation, laboratory waste disposal and many other support functions that allow the testing units to continue with their work.

## **QUALITY ASSURANCE AND SAFETY PROGRAM**

This section consists of a *Quality Assurance & Safety Officer* who is responsible for the quality assurance and safety programs within the Public Health Laboratories. The *Quality Assurance & Safety Officer* also directs the activities of the Quality Assurance Committee and the Safety and Emergency Response Committee (SERC). These committees are composed of volunteer staff members from every office and program located here at the Public Health Laboratories (PHL). This position is the reference person for safety and quality assurance related items.

### **PHL Quality Assurance Program**

The section coordinates the laboratory's compliance with all accreditation, proficiency and qualification regulations mandated by federal and state agencies, OSHA, EPA, HCFE, FDE, USDA, the Department of Energy and the Washington State Medical Test Site Rules. Additional QA functions performed by the Safety & QA Officer include:

- Coordinate the various subscribed or inter-laboratory proficiency testing programs.
- Maintain the quality assurance plan and consults with the laboratory's client groups
- Research and resolves client complaints
- Prepare for on-site inspections by internal or external groups that certify or accredit the Public Health Laboratory.
- Coordinate external College of American Pathologists, CAP, inspection of other laboratories per our CAP licensing requirement.
- Facilitate the performance of pipette, thermometer, and weights calibration checks.
- Recommend training as required for the facility.

### **PHL Safety Program**

The Safety & Quality Assurance Officer confers with and advises the laboratory director, managers, supervisors and employees on occupational safety and health issues. Plans, organizes and directs the laboratory's safety and health program to comply with OSHA, WISHA, IMR, the fire marshal and other applicable federal, state and local codes. Conducts accident investigations and inspections, recommends proper corrective or preventive actions. Additional safety functions performed by the Safety & QA Officer include:

- Collaborate with the Department of Health's risk management group, maintains, and updates the laboratory Chemical Hygiene Plan as required by WAC 296-62-400 and the other laboratory safety manuals and plans.

- Coordinate development of the Public Health Laboratory's disaster response plan, emergency response plan and evacuation plans/procedures in alignment with the Departmental plans.
- Investigate employee industrial and vehicular accidents.
- Coordinate claims and reports with the DOH risk manager.
- Conduct local facility/laboratory industrial safety inspections.
- Manages the Occupational Medicine program for the Public Health laboratory. Schedules immunizations, blood draws, etc.
- Conduct interviews with employees, supervisors and managers to identify/correct unsafe practices and conditions.
- Alternative Responsible Official for the Select Agent Program.
- Performs risk assessments to ensure that the appropriate control measures are implemented.
- Manages the Respirator Protection program. Performs respirator fit testing and training.
- Responsible for the management of the chemical waste handling and chemical inventory.
- Performs safety orientations for new employees with the employee's supervisor.
- Performs ergonomic assessments and works with the DOH Office of Risk Management to ensure that the PHL complies with WISHA regulations.
- Recommend safety related training.
- Review facility designs and make safety related recommendations.
- Reviews, with the *Safety and Emergency Response Committee*, the animal handling procedure for the facility.

## PUBLIC HEALTH LABORATORIES TRAINING PROGRAM

The PHL Training program has been conducting extensive laboratory training since it moved to the current facility in 1985. The facility includes a 1,035 square foot training laboratory complex, a classroom that will seat 24 people and a conference room capable of seating 90 people. Two full-time staff members are dedicated to provide training activities for both internal and external clients. Additionally, A bioterrorism training advisor has been added to the training program with funds provided by a grant from Centers for Disease Control and Prevention.

The PHL training staff develops and presents training courses for internal and external laboratory personnel. As a member of the National Laboratory Training Network operated by the Centers for Disease Control and Prevention and the Association of State and Territorial Public Health Laboratory Directors, the PHL Training program brings national training programs to Washington State.

### Training and Technical Assistance Provided

Conferences, symposia, workshops, seminars and bench training are scheduled for health care personnel throughout the state. A schedule of courses is posted on the web at <http://www.doh.wa.gov/EHSPHL/PHL/training/train.htm>.

For information on the Public Health Laboratories training and technical assistance call (206) 418-5402. Audio-visual materials are available upon request.

TRAINING PROGRAM				
PROGRAM SERVICES				
TRAINING	WHO CAN PARTICIPATE	SERVICES	WHEN	PHONE #
Audio-Visual Library	Anyone working in a clinical laboratory; teachers of laboratory science; health care providers.	Audio-visual training aids	Call to reserve	(206) 418-5404
Workshops Seminars Conferences	Announcements will describe target audiences	Given at the State Laboratories or in local facilities. Designed to meet current needs. May be lectures or lectures and wet workshops.	By announcement	(206) 418-5401
Bench Training	Working laboratory scientists with approval from lab director	New technology management training. Practical experience.	Call to arrange	(206) 418-5401
State Laboratories Tours, Public Relations and Support of Professional Organizations	Laboratory professionals, students, anyone with a special interest, health and laboratory groups	Opportunity to see a public health laboratory and understand how it serves the citizens of Washington.	On request Call to reserve	(206) 418-5401



TRAINING PROGRAM				
PROGRAM SERVICES				
TRAINING	WHO CAN PARTICIPATE	SERVICES	WHEN	PHONE #
Student Rotations or Internships	College students who have completed degree required microbiology, chemistry or health related coursework.	Opportunity to become familiar with public health careers in their chosen field. Practical experience.	Arrangements must be made through student's advisor.	(206) 418-5401
Post-Doc Rotations	Students who have completed course work for their doctorate degree.	This rotation will provide an opportunity to work on a public health project related to their degree.	Arrangements must be made through student's advisor.	(206) 418-5401



# OFFICE OF NEWBORN SCREENING

Washington State law (RCW 70.83 and WAC 246-650) requires that all babies born in Washington State be tested for certain conditions affecting newborns. The Office of Newborn Screening performs tests to detect the following nine treatable disorders:

- Biotinidase deficiency
- Congenital adrenal hyperplasia (CAH)
- Congenital hypothyroidism (CH)
- Cystic fibrosis (CF)
- Galactosemia
- Hemoglobinopathies (including sickle cell disease)
- Homocystinuria
- Maple syrup urine disease (MSUD)
- Medium chain acyl Co-A dehydrogenase deficiency (MCADD)
- Phenylketonuria (PKU)

Blood from a heel puncture is absorbed on a specialized filter paper; the card is then air dried and submitted to the unit *within 24 hours* for testing.

State law specifies that specimens from newborns must be collected prior to hospital discharge and no later than five days after birth. Specimens are to be submitted to the Office of Newborn Screening within 24 hours of collection. Parents may refuse testing on the basis of religious practices or tenets by signing a statement on the back of the NBS collection form. A fee is charged to parents through the hospital of birth. A second newborn screen is highly recommended at 7 to 14 days of age. There is no additional fee for these follow-up screening tests.

Health care providers may obtain screening kits from the Office of Newborn Screening, State Public Health Laboratories, 1610 N.E. 150th Street, Shoreline, WA 98155, phone (206) 418-5410 or toll free at 1-866-660-9050; fax (206) 418-5415.

Newborn screening specimens must be sent to:

Newborn Screening  
Washington State Department of Health  
PO Box 55729  
Shoreline, Washington 98155-0729

Place each specimen card in the protective envelope before mailing. Specimens *must not* be placed inside plastic bags, since this may cause the blood to degrade, thus invalidating the screening test results.

## Dried Blood Hemoglobin Testing

This unit provides testing of dried blood for hemoglobinopathies such as sickle cell disease for patients beyond the newborn period. These tests are available to aid health care providers in diagnosing hemoglobin disorders, and family members of affected individuals, and to screen high-risk groups. Hemoglobin testing is usually not indicated for children born in Washington after November 1, 1991, since these children were screened shortly after their birth as part of routine Newborn Screening.

### Turn-Around Times

Galactosemia.....1 day  
 Biotinidase, CAH, CH, Hemocystinuria, MSUD, MCADD, PKU, CF....2 days  
 Newborn Hemoglobin Screening .....5 days  
 Dried Blood Hemoglobin Testing .....11 days

NEWBORN SCREENING						
Collection and Submission Instructions						
AGENT/ DISEASE	SPECIMEN	COLLECTION TIME	CONTAINER	TRANSPORT TEMP.	STORAGE TEMP.	REMARKS
Congenital hypothyroidism, phenylketonuria, congenital adrenal hyperplasia & hemoglobinopathies (required by law on every newborn in the state) biotinidase deficiency, galactosemia, homocystinuria, maple syrup urine disease (MSUD) and medium chain acyl Co-A dehydrogenase deficiency (MCADD) will be added by June 2004.	Blood from heel stick saturated on filter paper and dried at ambient temperature	Prior to hospital discharge, no later than five days after birth. A second specimen is recommended at 7-14 days.	Specialized filter paper included on specimen collection forms	Air dry at ambient temperature (2 hours minimum); place into individual protective envelopes: mail within 24 hours of collection (ambient temperature)	Ambient temperature; ship within 24 hours of collection	For additional information & educational materials, call (206) 418-5410 or toll free at 1-866-660-9050  On the web: <a href="http://www.doh.wa.gov/nbs">www.doh.wa.gov/nbs</a>
Hemoglobin conditions including sickle cell disease and trait (outside of the newborn period)	Blood saturated on filter paper and dried at ambient temperature	Any	Specially modified specimen/filter paper card (similar to above).	Same as above.	Same as above.	For additional information, call (206) 418-5410 or toll free at 1-866-660-9050  On the web: <a href="http://www.doh.wa.gov/nbs">www.doh.wa.gov/nbs</a>

## OFFICE OF PUBLIC HEALTH MICROBIOLOGY

This office provides consultation and training to other laboratories, hospitals, health care providers and local health/environmental jurisdictions to enhance technical skills, productivity, efficiency, and to assure quality service. It carries out a wide range of microbiology surveillance activities, including isolation, definitive antimicrobial identification, molecular diagnostics, drug sensitivity and/or confirmation of etiological agents of public health and epidemiological concerns.

### ENTERIC BACTERIOLOGY

The Enteric Laboratory serves primarily as a reference laboratory for the identification of enteric pathogens. The unit also serves Local Health Jurisdictions by screening clinical samples for enteric pathogens; i.e., *Salmonella*, *Shigella* and *Campylobacter*. Examinations for *Yersinia*, *Vibrio* and *E. coli* O157:H7 are available to local health jurisdictions/districts upon request.

Washington State Board of Health Regulations (WAC 246-100) require that all *Salmonellosis* (including typhoid fever), *Shigellosis* and *V. cholerae* isolates be confirmed by the State Enteric Laboratory. All *E. coli* O157:H7 isolates submitted are subtyped using Pulsed Field Gel Electrophoresis (PFGE). All laboratories are requested to submit isolates of *E. coli* O157:H7 for PFGE testing.

#### Turn-Around Times

##### *Salmonella* spp.

Presumptive identification.....	3 days
Final identification .....	10 days

##### *Shigella* spp.

Presumptive identification.....	3 days
Final identification .....	4 days

##### *E. coli* O157:H7

Presumptive result .....	3 days
Final identification .....	6 days

##### *Campylobacter jejuni*

Final identification .....	6 days
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##### *Vibrio*

Presumptive identification.....	4 days
Final identification .....	5 days

##### *Yersinia enterocolitica*

Reference culture	
Presumptive identification.....	3 days
Final identification .....	5 days

##### Stool

Presumptive identification.....	3 days
Final identification .....	up to 3 weeks

<b>ENTERIC BACTERIOLOGY</b>				
<b>Collection and Submission Instructions</b>				
<b>AGENT / DISEASE</b>	<b>SPECIMEN &amp; QUANTITY</b>	<b>COLLECTION TIME</b>	<b>CONTAINER</b>	<b>REMARKS</b>
Reference Cultures (Isolates Only)	Send only pure cultures		Double Mailer	Non-Fermentative agar media; screw-cap tube; tighten cap; *use Pai or Loeffler agar for <i>Salmonella typhi</i> ; <i>Campylobacter</i> use blood agar slant for submission of isolates for confirmation.
<i>Salmonella</i> (not Typhoid)	Swab coated with stool specimen or rectal swab	At onset	Cary-Blair transport media, double mailer	
<i>Salmonella typhi</i>	Stool: Swab coated with stool specimen or rectal swab	Stool: 2 - 3 days after onset	Stool: Cary-Blair transport media, double mailer	
	Urine: Volume equal to amount of Buffered Glycerol Transport		Urine: Request Buffered Glycerol Saline transport media	Urine: Buffered Glycerol Saline transport media is available by calling the Enteric Unit (206) 418-5456.
<i>Shigella</i>	Swab coated with stool specimen or rectal swab	At onset	Cary-Blair transport media, double mailer	
<i>Vibrio</i>	Swab coated with stool specimen or rectal swab	At onset	Cary-Blair transport media, double mailer	Call the Enteric Unit (206) 418-5456 before sending. Specify test on Enteric form.
<i>Campylobacter</i>	Swab coated with stool specimen or rectal swab	At onset	Cary-Blair transport media, double mailer	Use blood agar slant for submission of isolates for confirmation.
<i>Yersinia enterocolitica</i>	Swab coated with stool specimen or rectal swab	At onset	Cary-Blair transport media, double mailer	Test requires three (3) weeks; specify test on Enteric form.
<i>Escherichia coli</i> O157:H7	Swab coated with stool specimen or rectal swab	At onset	Cary-Blair transport media, double mailer	Specify test on Enteric form; PFGE testing included; CDC performs tests for SLT I & II on non-motile isolates
Shiga toxins I, II (verotoxins) produced by enterohemorrhagic <i>E. coli</i>	Swab coated with stool specimen or rectal swab	At onset	Cary-Blair transport media, double mailer	EIA for detection of Shiga toxins I, II (verotoxins) available – please specify on Enteric form.

\* NOTE: All culture samples (except *Salmonella typhi*) use double mailers and slant screw-cap tubes with non-fermentative media.

## MOLECULAR DIAGNOSTICS

Working together with the Molecular Epidemiologist on staff, the Microbiology laboratories have been able to collaborate with local, national and international universities, the Centers for Disease Control and Prevention (CDC) in Atlanta, GA, as well as other state public health laboratories to implement new types of DNA based technology having an application to public health. The new technologies currently in development and use include PFGE, PCR, fluorescent *in-situ* hybridization (FISH), TaqMan (a real-time PCR) and DNA sequencing. All PCR submission requests must be pre-approved by your Local Health Jurisdiction and Dept. of Health Communicable Disease Epidemiology.

### Polymerase Chain Reaction (PCR) Unit

SERVICES OF POLYMERASE CHAIN REACTION (PCR) UNIT				
Agent/Disease	Specimen	Collection Time	Turn-Around Time	Remarks
Diarrhea due to Norovirus	Stool	Illness	Tests are performed as requested	RNA extraction followed real-time PCR
Encephalitis due to West Nile Virus	CSF or haemolymph of the mosquito	Illness	Tests are performed as requested	Taqman PCR
Whooping cough due to B. pertussis	Dacron Nasopharyngeal swab	Illness	Tests are performed two times per week. *	DNA extraction followed by real-time PCR PCR testing must be pre-approved by the County Health Dept.
Legionella longbeachae and pneumophila	Isolate plated out on BCYE agar		Tests are performed three times per week. *	Real-time PCR and Fluorescent in situ hybridization (FISH)
SARS	Stool or respiratory samples		Tests are performed as requested	RNA extraction followed real-time PCR
Influenza A	NP swabs or respiratory washes		Tests are performed as requested	RNA extraction followed real-time PCR
Mumps	Serum		Tests are performed as requested	RNA extraction followed real-time PCR

\* Test will be performed immediately if so requested by a State Epidemiologist.

### Pulsed Field Gel Electrophoresis (PFGE) Unit

PFGE testing is used to genetically compare two or more strains of bacteria. *All specimens submitted to the PFGE Unit should be pure bacterial isolates.*

#### Turn-Around Times

All pure bacterial isolates other than Mycobacteria\*:..... 2 days

\* The Mycobacteriology isolates will be sent to a CDC genotyping lab for genotyping

PFGE TESTING SERVICES	
<b>BACTERIAL DISEASES/AGENTS</b>	Any bacteria other than <i>Mycobacteria</i> *
<b>ACCEPTABLE CLINICAL CULTURES</b>	Pure isolated cultures only
<b>ID OR CONFIRMATION</b>	No
<b>TESTS FOR TOXINS/TOXICITY</b>	No
<b>YOUR ID &amp; RESULTS REQUIRED</b>	Yes
<b>DNA TYPING</b>	Yes
<b>CALL BEFORE SHIPPING</b>	Yes, Call: (206) 418-5561
<b>CONFIRMATION REQUIRED BY STATE LAW</b>	
* The Mycobacteriology isolates will be sent to a CDC genotyping lab for genotyping	

## MYCOBACTERIOLOGY

The Mycobacteriology Unit serves as a reference laboratory for the identification of mycobacteria. This unit also offers isolation and identification of mycobacteria from clinical specimens.

Susceptibility testing for the first and second line anti-tuberculosis drugs is performed in this unit on isolates of *M. tuberculosis* complex (Mtb). For a more extensive drug susceptibility profile, isolates are sent to Centers for Disease Control and Prevention (CDC). It can take longer for drug susceptibility results if the patient shows resistance to any drugs. Submitters requesting results faxed to them must submit a *CONFIDENTIALITY NOTICE* stating that their fax machine is in a secure location accessible *ONLY* to authorized personnel.

### Turn-Around Times

#### Smear Results:

Positive smear (*phoned, mailed and faxed*). ..... 24 - 48 hours  
Negative smear (*mailed*). ..... 24 - 48 hours

#### Amplified Mycobacterium Tuberculosis Direct (AMTD) Test:

*Performed on first-time positive smears and on special requests with prior arrangement*  
Results (*phoned and faxed upon request*) ..... 24 hours

#### Culture Results:

AFB positive cultures  
(*phoned, mailed and faxed upon request*) ..... 1 - 4 weeks  
AFB Negative cultures (*mailed*). ..... 8 weeks

#### Genetic Probe Test (GP):

*Performed weekly on AFB positive cultures*  
Results (*phoned and mailed*)..... 1 - 4 weeks

If genetic probes are negative for Mtb, *M. avium* complex (MAC), *M. goodii*, and *M. kansasii*, then the culture is considered *atypical* or a MOTT (Mycobacterium Other Than Tuberculosis), and will be identified using biochemical analysis.

#### Biochemical Analysis

Results (*phoned and mailed*) ..... 4 months

### Drug Susceptibility Test

*Performed on confirmed Mtb cultures*

Bactec\* ..... 7 - 10 days  
Plate method\*\* ..... 4 weeks

\* Bactec drugs employed: Streptomycin, Isoniazid, Rifampin, Ethambutol, Pyrazinamide.

\*\* Plate method drugs employed: Streptomycin, Isoniazid, Rifampin, Ethambutol, Ethionamide, P-aminosalicylic acid, Ofloxacin, Amikacin.



MYCOBACTERIOLOGY					
Collection and Submission Instructions					
AGENT/ DISEASE	SPECIMEN	COLLECTION TIME	CONTAINER	TRANSPORT TEMP.	REMARKS
Mycobacteria	Sputum	Early morning specimen on 3 consecutive days, ship each specimen as obtained to be received by PHL less than 5 days after collection	Sterile plastic centrifuge tube**	Ambient temperature	Two to three teaspoonfuls are sufficient. Saliva is a poor specimen. Send each specimen as collected. *
	Gastric washing	Before breakfast, early morning specimen on 3 consecutive days are recommended	Sterile plastic centrifuge tube**	Ambient temperature or Refrigerate at 4°C	Send each specimen as collected*
	Urine	Early morning midstream collection on 3 consecutive days, 30 ml per tube, ship each specimen as it is obtained	Sterile plastic centrifuge tube**	4°C, if possible	Send each specimen as collected, tighten cap well and seal pressure-sensitive labeling tape*
	Stool	See remarks	Sterile specimen container or centrifuge tube	Ambient temperature	Specimens must be received at the TB Unit within 24 hours of collection, call TB Unit before shipping*
	Spinal fluid		Small, tightly capped, sterile containers	Ambient temperature or 4°C	Ship as indicated*
	Tissues or swabs	Collect aseptically	Same as above	Refrigerate at 4°C	Should have a small amount of sterile distilled water added to prevent drying. * Carey Blair or Aimes transport media NOT recommended.
Mycobacterial cultures	For ID, confirmation or susceptibility testing		Culture tube, securely tighten cap, seal pressure-sensitive labeling tape*	Ambient temperature	Ship as indicated*, use courier service to ship Petri dishes
* Specimens must be shipped in double cardboard mailers to meet IATA, OSHA and postal requirements. ** Specimen collection kit may be ordered through PHL.					
<b>Pursuant to WAC 246-101, positive results for <i>Mycobacterium tuberculosis</i> are notifiable within 2 working days to DOH – TB Laboratory. Specimen submission is required.</b>					

## SEXUALLY-TRANSMITTED DISEASES

The Sexually-transmitted Diseases Unit functions primarily as the reference laboratory for the state for the definitive identification or confirmation of cultures, which have been isolated and tested in other laboratories. Reference specimens are accepted from all laboratories. Clinical specimens are only accepted from local health jurisdictions STD clinics. Call (206) 418-5492.

Chlamydia and *N. gonorrhea* clinical specimens are tested by the Gen-Probe Aptima II Combo test. This is a genetic amplification test and allows the testing of both Chlamydia and *N. gonorrhea*. Clinical specimens are accepted from local health jurisdictions STD clinics, Family Planning Clinics and Planned Parenthood Clinics.

### Turn-Around Times

*N. gonorrhea* clinical specimen ..... 1 - 2 days  
*N. gonorrhea* reference culture ..... 2 - 5 days  
 Chlamydia/GC..... 2 - 4 days

SEXUALLY-TRANSMITTED DISEASES			
Collection and Submission Instructions			
AGENT/DISEASE	SPECIMEN	COLLECTION	REMARKS
Gonorrhea Clinical specimen	Culture of urethral exudate, cervix, rectum and throat	Illness or contact	Follow special direction with kits, temperature for incubation and media used are important
Reference culture	All sites		Submit in screw-capped tube, sealed pressure-sensitive labeling tape. Use chocolate agar slant or modified Martin-Lewis pill pocket plate and place in CO <sub>2</sub> biohazard bag. Incubate 35° C 24 hours before shipping.
Chlamydia/ <i>N. gonorrhea</i>	Swab of cervix	Project criteria, use collection kits from PHL	This is a Federal project and only participants may send specimens

## SPECIAL PATHOGENS SURVEILLANCE (REFERENCE)

The Special Bacteriological Pathogens Unit functions primarily as a reference laboratory for confirmation or definitive identification of cultures isolated and tested in other laboratories. The unit uses the Hewlett Packard Microbial Identification System (MIS) along with a variety of biochemical and serologic methods to identify these organisms. The selection and extent of the tests used for identification vary according to the origin of the specimen from which the microorganism was isolated and the type of infection suspected or produced. This information must be provided on the Reference Bacteriology Laboratory Request Form before processing can begin. Clinical, Environmental and Bioterrorism specimens may be accepted for organisms, which produce infections such as anthrax, human brucellosis, botulism (food, wound, infant), *cholera* and non-*cholera* Vibrios, glanders, melioidosis, plague, tetanus, tularemia, relapsing fever and Legionnaires' disease.

Unless there is a special problem, the unit does not test animal and environmental isolates (if there is an suspect/ identified public health problem, the Special Pathogens lab will test with the approval of Dept. of Health's Epidemiology Section (206-418-5500) or toll free (1-877-539-4344).

Every effort should be made to send a pure, viable culture. Each submitter should maintain a subculture of the organism submitted until the final identification is received.

The unit requests that records of the existing work performed on each culture be included when submitting the culture. The suggested laboratory results accompanying each specimen are listed in the table below.

SPECIAL PATHOGENS SURVEILLANCE (Reference)	
Required Test Results	
<b>Aerobes:</b>	Gram stain reaction Colony morphology Catalase Oxidase Motility Carbohydrate reactions Temperature studies (when pertinent) Spore formation (when pertinent)
<b>Anaerobes:</b>	Gram stain reaction Colony morphology Catalase Motility Spore formation (when pertinent)

Most aerobes can be mailed on blood, chocolate, nutrient, trypticase soy or brain heart infusion agar slants. Anaerobe cultures can be successfully sent in screw- capped tubes of chopped meats, brain broth, fresh thioglycollate broth, boiled motility media, or a commercial transport system. Before shipping anaerobes, overlay the media with  $\frac{3}{4}$  inch

of sterile Vaspar, petrolatum or paraffin. Tape the tightened caps with tape. **DO NOT MAIL PETRI DISHES.**

### **Turn-Around Times**

Reference Cultures .....7 - 60 days

### **Bioterrorism Response Laboratory**

The Special Pathogens Unit provides laboratory support for bioterrorism response to requests from local Health Jurisdictions, Federal Bureau of Investigation, U.S military and law enforcement. All requests for laboratory bioterrorism response must pass through the local health dept. who will alert the State Health Dept. Epidemiology Unit (206-418-5500) or toll-free 1-877-539-4344. Environmental (powders, liquids, etc.) specimens for Bioterrorism response should first involve local law enforcement. Bioterrorism response is available around-the-clock. The Special Pathogens Unit does not accept bioterrorism samples from the public or from commercial entities.

Clinical samples and bacterial isolates can be sent to the Special Pathogens lab. Submitting institutions should first call the local Health Dept. See **Appendix B – Collection Instructions for Terrorism-Related Samples.**

### **Special Bacteriology Requests Turn-Around Times**

#### ***Bacillus anthracis* (anthrax)**

Presumptive culture.....1 - 4 hours

Confirmed .....48 hours

#### ***Clostridium botulinum* (botulism)**

Presumptive Toxin .....2 - 6 hours

Confirmed Toxin.....4 days

Culture.....7 - 14 days

#### ***Francisella tularensis* (tularemia)**

Presumptive Culture confirmation .....1 - 4 hours

Culture Isolation.....2 - 7 days

#### ***Yersinia pestis* (plague)**

Presumptive.....1 - 4 hours

Confirmed .....14 days

#### ***Brucella* species**

Culture.....10 - 31 days

#### ***Legionella* species**

DFA.....1 – 4 hours

Culture.....3 - 7 days

Environmental (pre-approved by epidemiology).....10 - 14 days

#### ***Burkholderia pseudomallei***

Presumptive.....24 hours

Final.....7 days

#### ***Burkholderia mallei***

Presumptive.....24 hours

Final.....7 days

Ricin toxin.....8 hours

(CDC confirmation 3 days)

## **Stock Cultures**

The Special Bacteriological Pathogens Unit has a collection of reference bacteria. These organisms are available to any laboratory in the State upon request. For reasons of time and expense involved, the laboratory must limit the number of requests from any one laboratory. Included in our collection are a variety of less commonly encountered organisms. The Special Pathogens Unit does not send out stock cultures that pose a significant public health threat. These include *Bacillus anthracis*, *Brucella* spp., *Yersinia pestis*, *Francisella tularensis*, *Burkholderia pseudomallei*, *Burkholderia mallei*, *Clostridium botulium*, and *Salmonella typhi*.

Please consult with the unit before ordering stock cultures (206) 418-5452. They can also discuss with you the best way to maintain the requested organisms.

Requests for cultures must be made on your official letterhead.

## Services of Special Pathogens Surveillance

Bacterial Diseases/Agents	Acceptable Clinical Cultures	Id or Confirmation	Tests For Toxins/ Toxicity	Antimicrobial Susceptibility Tests	Your Id & Results Required	Call Before Shipping (206)418-5452	Typing	Confirmation Required By State Law
Anaerobic infections due to Bacteroides								
Anthrax				*				
Botulism (Food, Wound, Infant)								
Brucellosis (human)								
Cholera and non-cholera Vibrios						**		
Clostridia								
Cocci (anaerobic)								
Corynebacterium spp.								
C. diphtheriae								
"Enteric" (Enterobacteriaceae) infections				*				
Erysipeloid								
Fusobacterium								
Legionella pneumophila infections								
Listeriosis								
Melioidosis								
Neisseria meningitidis Spinal Fluid (CSF) & Blood Isolates								
Miscellaneous Gram "-" rod								
Miscellaneous Gram "+" rod								
Plague				*				
Pneumococcal infection				*				
Rat bite fever								

## Services of Special Pathogens Surveillance

Bacterial Diseases/Agents	Acceptable Clinical Cultures	Id or Confirmation	Tests For Toxins/ Toxicity	Antimicrobial Susceptibility Tests	Your Id & Results Required	Call Before Shipping (206)418-5452	Typing	Confirmation Required By State Law
Relapsing fever								
Staphylococcal infections								
Streptococcal infections								
Tetanus								
Tularemia				*				

## Special Pathogens Collection and Submission Instructions

<i>Clostridium botulinum</i>				
Type Of Botulism	Specimen	Collection	Results/Tat	Remarks
Food Botulism	Serum	A 5 - 15 ml specimen (preferred) collected soon after onset of symptoms and before antitoxin is given	All results are from 4 hours to 14 working days for Food and Infant & wound botulism	Advise Lab if any drugs have been given Specimens (food, stool, serum) should be submitted on suspect cases. See WAC 246.100-231 for further details
	Gastric Material	Walnut-size (50 gm)		For all specimens, unless otherwise specified:
	Stool	10 - 50 grams (preferably walnut-size). Enema material is acceptable. Obtain specimen from sterile (non-bacteriostatic) water or saline enema. A volume of 20 ml collected after enema is sufficient.		Place specimen in a sterile, leak-proof container, place in plastic bag, then in an insulated shipping container with ice packs
	Vomit	10 - 15 ml		Ship cold*
	Food	Unopened food, food remnants, dishwasher-washed/unwashed container		DO NOT FREEZE
Infant Botulism	Stool	Frequently difficult to obtain a sufficient quantity. Obtain specimen from sterile (non-bacteriostatic) water or saline enema. A volume of 20-30 ml collected is sufficient.		

	Autopsy Specimens	Intestinal samples should be taken from different levels (small bowel, proximal colon, distal colon)		Prior approval by CD Epidemiology required (206) 418-5500
Wound Botulism	Serum	Same as serum above	4-96 hours	Notify Special Bacteriological Pathogens Unit as to when and how specimens are being shipped (206) 418-5452
	Tissue	Representative tissue sample	2-14 working days	
	Swab	Swab place in anaerobic transport media	2-4 working days	Ship ambient temperature
<b>Legionella (Legionnaire's Disease) Culture and DFA</b>				
<b>Agent/ Disease</b>	<b>Specimen</b>	<b>Collection</b>	<b>Transport Temp.</b>	<b>Remarks</b>
Culture	Bronchial washing, bronchoalveolar lavage, pleural fluid, sputum or trans-tracheal aspirates, fresh lung biopsy material	Illness	Keep cool with ice packs	Sterile, screw-capped container or tube
DFA Slides	Preserve specimens in 10% formalin or slides cut from paraffin sections	Submit a minimum of 5 slides with 2 test areas	Ambient temperature	With all Legionella specimens
Tissue	Preserve specimens in 10% formalin or slides cut from paraffin sections	Submit a minimum of 5 slides with 2 test areas	Ambient temperature	Ship promptly  Transport slides in slide carriers
Environmental Prior approval by State Epidemiology required	1-10 liters water, swabs from faucet, cooling tower water, soil	Patient with symptoms compatible with legionnaires' Disease	Keep cool with ice packs	Ship promptly, Specimen collection usually performed by State Lab personnel

Services are provided for the diagnosis and confirmation of those infectious disease agents that are of a public health nature. Group A *Streptococcus* isolates from patients who present diagnostic problems and require reference services will be accepted. Local laboratories should be utilized for routine testing for Group A *Streptococcus*. Clinical specimens for *Corynebacterium diphtheria* and *Bordetella pertussis* are accepted directly from laboratories and local health jurisdictions.

### Turn-Around Times

#### *Bordetella pertussis*

Culture.....7 - 10 days  
DFA.....1 - 2 days  
PCR.....2 – 3 days

#### *Corynebacterium diphtheriae*

Clinical .....2 - 3 days  
Culture.....2 days  
Toxin Test .....4 days



*Corynebacterium ulcerans*

Clinical .....	2-3 days
Culture.....	2 days
Toxin Test .....	4 days

*Group A streptococci*

Clinical/Culture .....	1 - 2 days
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<b>SPECIAL RESPIRATORY PATHOGENS</b>			
Collection and Submission Instructions			
Agent/Disease	Specimen	Collection Time	Remarks
Bordetella pertussis Clinical Culture  DFA  PCR	Swab left and right nasopharyngeal areas  Swab left and right nasopharyngeal areas  Dacron Nasopharyngeal swab (left and right areas)	Illness  Illness Illness	Streak one set of swabs of on Kendrick-Jones charcoal agar, leave swabs in tube  Streak another set on two microscope slides, air dry Tests are performed two times per week.* PCR testing must be pre-approved by the County Health Dept.
Corynebacterium diphtheria Throat, N/P  Wound Culture  Reference Culture	Swab both of the nares to the posterior pharyngeal wall and the oral pharynx  Clean wound site with sterile, normal saline removing crusted material  Submit on Pai slant, Loeffler slant, or blood agar slant	Illness, contact or carrier	Obtain nasopharyngeal cultures with a flexible alginate swab, take throat cultures with a cotton - or Dacron swab which is firmly applied to any area with a membrane or inflammation, streak nasal specimen on one Pai slant; the oropharynx on the other pai slant, leave swabs in tube.  Place cotton - or type swab firmly to base of the wound, streak swab on Pai slant, leave swab in tube
Group A streptococci Clinical  Reference Culture	Tonsils and pharynx should be rubbed with a cotton- or Dacron-tipped swab, touch any exudate with the swab, avoid the tongue and uvula tissues	Illness	Streak one swab on Pai slant, leave swab in tube, place second swab in silica gel tube  Submit on blood agar slant

\* Test will be performed immediately if so requested by a State Epidemiologist.

## SYPHILIS SEROLOGY

The Syphilis Serology Unit serves primarily as the reference laboratory for the confirmation of sera results that are reactive by any serological test for syphilis. The Venereal Disease Research Laboratory (VDRL) test is performed on all sera and spinal fluids submitted to the Syphilis Serology Unit. If the result is weakly reactive (WR) or (R) a confirmatory test that is specific for *Treponema pallidum* antibody (TP-PA) is performed.

The TP-PA is not routinely performed on sera that are non-reactive. Exceptions can be made but must be communicated to the Serology Unit by checking the Reference box on the requisition slip.

The VDRL is used to evaluate the results of treatment therapy as it tends to revert to a lower titer or non-reactive after treatment. The TP-PA will remain positive after treatment.

### Test Interpretation - Serum and Spinal Fluid

VDRL results	TP-PA results*:
Non-reactive (-)	Non-reactive (-)
Weakly reactive, Titer of 1:0	Reactive (+)
Reactive, Titer of 1:1 or greater	

\*Spinal fluid rarely produces biological false positive reactions. A reactive spinal fluid usually indicates tertiary syphilis.

### Turn-Around Times

VDRL.....	5 days
TP-PA.....	6 days

SYPHILIS SEROLOGY					
Collection and Submission Instructions					
AGENT/ DISEASE	SPECIMEN	COLLECTION	TRANSPORT TEMP.	STORAGE TEMP.	REMARKS
Syphilis	Draw 5 - 10 ml sterile whole blood in a tiger- or red-top tube	Diagnosis or treatment follow- up	Refrigerated is preferred, ambient temperature transport is acceptable	Refrigerator (4°)	Use sterile chemically clean tubes, syringes, etc.  Never freeze whole blood.
	2.0 ml serum	Same		-20°C freezer or refrigerator	Do not send plasma.  Transport promptly
	0.5 ml Cerebral Spinal Fluid (CSF)	Diagnosis or treatment follow- up			

## Premarital Blood Testing

Washington State law does not require a premarital blood test for syphilis; however, some states do. Premarital testing can be performed by any laboratory which is a Medical Test Site licensed and certified for Syphilis Serology. The laboratory performing the test should obtain and fill out the premarital certificate that is then submitted to the patient's doctor for signature. The Public Health Laboratories have Premarital Certificates for those states requiring premarital Syphilis Serology testing and can perform a test for syphilis.

## Information Required on Lab Form for Premarital Certificate

Patient's *full name including middle name spelled out*, patient's address, age, sex, date blood drawn, name and address of doctor and state where patient is getting married. This information is required on the premarital certificate.

## Testing Information

A doctor licensed to practice in Washington State must request syphilis blood tests. A time limit is stated on the premarital certificate.

***Deliver blood and laboratory form to Washington State Public Health Laboratories (PHL) by courier, mail or by patient.*** Non-reactive results can be provided within two workdays of receipt. Same-day results can be provided ***only if*** patient calls PHL at (206) 418-5444, delivers blood, and picks up premarital certificate at PHL. ***Reactive results must be confirmed, which requires an additional three workdays.***

After the test is completed, the doctor who requested the test must sign the certificate.

Some states also require a rubella test for women of childbearing age. This test is not available at the State Laboratories, but most private laboratories do this test. The Seattle-King County Laboratory also does rubella testing.

States Requiring Syphilis	States Requiring Rubella
Connecticut	Connecticut
Georgia	Georgia
Massachusetts	Indiana
Mississippi	Massachusetts
New Jersey	Montana
New Mexico	Nebraska
Oklahoma	New Mexico
Pennsylvania	Rhode Island
Rhode Island	
West Virginia	
District of Columbia	<u>HIV Testing</u>
<u>Other Areas—</u>	Rhode Island requires HIV test unless waiver is signed
Puerto Rico - syphilis only	
Mexico – syphilis	

## VIROLOGY

The Virology Unit provides reference, surveillance and diagnostic services. Laboratory diagnosis of viral agents can be done by culturing for the virus, serologic testing for associated antibody, or by antigen detection. The preferred methodology varies with each virus; see the following tables for information about specific viruses. Ship all specimens cold, labeled with patient's name, for overnight delivery. Specimens without a patient name written on the tube may not be tested. A "Virus Examinations" lab form, with patient name, type of specimen, date of onset, date of collection, test requested, and submitter name and address must accompany all specimens.

Most viral testing requires approval by Epidemiology (phone 206-418-5500) before sending the specimens. These tests are for detection of influenza, measles, Norovirus, rabies, rubella, SARS, St. Louis Encephalitis, and West Nile virus.

VIROLOGY	
Testing Information	
<b>Culture</b>	<i>Culture, or isolation, of virus is performed by inoculating the specimen into tissue culture cells. For best recoverability and survival of a virus, 1) collect within three days of onset of symptoms, 2) place swabs in Viral Transport Medium (VTM), and 3) keep the specimen cold at all times, including during shipment.</i>
<b>Serologic Testing</b>	<i>All serology tests require prior approval by Epidemiology. Testing of serum (or limited testing of CSF) by enzyme-linked immunoassay (ELISA) detects IgA, IgM, and IgG antibodies to determine recent infection. The PHL performs ELISA only on people who are sick. Collect blood in a red top or red-gray top tube at the following times: measles, 3 days post onset of rash; rubella, 5 days post onset of rash; SARS, 21 days or more post onset of symptoms; West Nile and SLE, 8 days post onset of symptoms. When blood is drawn before these times, a second draw is sometimes needed to verify the results. For SARS testing, other specimens can be collected at earlier times as specified by Epidemiology.</i>
<b>Antigen</b>	<i>Directigen, Direct Fluorescent Antibody (DFA), and real time reverse transcription polymerase chain reaction (RT-PCR) can detect viral antigen in the original specimen. Directigen Flu A is performed on respiratory specimens collected by selected nursing homes and physicians. DFA can detect Varicella-Zoster virus in vesicular fluid. DFA is also done on animal brain to detect rabies. RT-PCR can detect antigen from West Nile virus and SARS in various specimens.</i>
<b>Rabies</b>	<i>To determine when an animal needs to be tested for rabies, consult your local health jurisdiction (LHJ), which will coordinate submission of the animal brain to the Virology unit. A DFA test detects rabies virus. All results of rabies testing are reported by telephone, in addition to mail, to the submitting LHJ. Emergency (evenings and weekends) rabies testing can be arranged by consultation with Epidemiologist on call, reached by calling (206) 418-5500.</i>
<b>PCR</b>	<i>Influenza A &amp; B can be identified by real-time reverse transcription PCR. Influenza A can be subtyped to H1, H3, H5 or H7. Either sputum, respiration or secretions, or nasal pharyngeal swabs can be submitted. Dacron swabs may be shipped dry if only PCR is requested. If both PCR and culture are requested, place Dacron swab in VTM.</i>

Virus Testing			
Symptoms and Virus	Specimens for Isolation	Type of Serology	Other Tests
<b>Respiratory Symptoms</b>			
Adenovirus Enterovirus HSV Influenza Virus Mumps Virus Parainfluenza Virus RSV SARS	NW, NP/THR NW, NP/THR NW, NP/THR NW, NP/THR Parotid gland, Urine NW, NP/THR NW, NP	ELISA – IgA/ IgG/ IgM	Directigen Flu A PCR  SARS RT-PCR
<b>Rash Symptoms</b>			
<b>Vesicular</b> Enterovirus HSV Varicella-Zoster (VZV)  <b>Maculopapular</b> Adenovirus  Measles (Rubeola) Rubella virus	NW, NP/THR, Stool, VF VF VF  NW, NP/THR, Stool	ELISA – IgG, IgM ELISA – IgG, IgM	VZV DFA, PCR
<b>CNS Symptoms (meningitis, encephalitis)</b>			
Adenovirus Enterovirus HSV Mumps  Rabies  St. Louis Encephalitis (SLE) West Nile virus (WN)	NW, NP/THR, CSF NW, NP/THR, Stool, CSF CSF, Brain NW, NP/THR, Urine, CSF	ELISA – IgM, IgG ELISA – IgM, IgG	Rabies DFA  WN RT-PCR
<b>Congenital, Perinatal Symptoms</b>			
Enterovirus HSV	NW, NP, THR, Stool, CSF NW, NP/THR, VF		
<b>Gastrointestinal Symptoms</b>			
Adenovirus Enterovirus Norovirus	Stool Stool Stool		
<b>Genital Lesions</b>			
HSV	VF, ES		

**Legend:** CSF: cerebral spinal fluid, DFA: direct fluorescent antibody, ES: endocervical swab IgA: initial antibody formed, transient (half-life of 4-5 days), IgG: antibody formed 2-3 weeks after onset of disease, IgM: antibody formed soon after onset of symptoms, between IgA and IgG, NP: nasopharyngeal, NW: nasal wash, RT-PCR: reverse transcription polymerase chain reaction, THR: throat, VF: vesicular fluid.

## Specimen Guide Notes:

1. The onset date of symptoms is needed for ALL SPECIMENS.
2. Send ALL SPECIMENS cold.
3. Isolation, DFA, and PCR:
  - a. Ideally, collect specimen within three (3) days of the onset of symptoms. More than 3 days will result in a lower virus load and may cause a failure to detect virus.
  - b. Send cold within 24 hours of collection for overnight delivery. Varicella-Zoster virus and measles virus in particular are fragile viruses.
4. Serology:
  - a. Specimens for measles antibody testing may be positive by the first day of rash; 95% of measles cases are positive by the third day after rash onset.
  - b. Rubella cases are slower to form antibody than measles cases. If the initial rubella IgM test is negative and the clinical diagnosis of rubella has not been ruled out, draw another serum seven (7) days after the first serum.
  - c. For other serology tests, an acute serum is one that is drawn within 7 days of the onset of the symptoms. A convalescent serum is a serum drawn 2 – 3 weeks after the acute. There is an advantage to draw both. In positive cases, a rising antibody level indicates recent illness; a stable antibody level indicates illness at some time in the past.
  - d. For arbovirus serologies, travel history one month prior to the onset and a history of insect bites during the travel should be entered on the lab form. Acute and convalescent sera are strongly preferred.

**Note:** This specimen guide is only a partial list of viruses that may be tested by reference laboratories. Call the Virology laboratory to ask about a particular viral test. If the specimen will be tested by Centers for Disease Control, a two page (8 1/2 x 11 size) form needs to be filled out and can be faxed, upon request.

### Turn-Around-Times

Virus Culture .....	7-14 days
Virus Serology .....	3–7 days
ELISA IgA, IgM, IgG .....	1-3 days
Antigen Detection.....	10 days
Rabies .....	1 day
PCR .....	1 day

### HIV/AIDS (Acquired Immune Deficiency Syndrome)

This unit provides HIV clinical laboratory serum testing ONLY to local health jurisdictions and contracted sites approved by the HIV program. The Request for Antibodies to HTLV III form must be completed for testing. Identify the specimen by patient's initials, birth date and/or a patient number.

The causative agent for Acquired Immune Deficiency Syndrome (AIDS) is the Human Immunodeficiency Virus (HIV). The previous name was HTLV III/LAV.

The screening procedure for serum specimens is an ELISA or EIA test. If the ELISA or EIA is a repeatable reactive, (reactive in two separate runs) a supplemental test is done. The supplemental test is the Western Blot.

**HIV Turn-Around Times**

ELISA/EIA..... 1 - 2 days  
Western Blot.....5 days



## **Appendix A: Shipping Information For PHL Clients**

### ICAO Guidance Document

#### Packaging and Labeling Checklists:

- Method of Transport
- Infectious Substance: Surface
- Infectious Substance: Transport via Air
- Infectious Substance: Post Office
- Diagnostic Specimens: Transport via Surface
- Diagnostic Specimens, Risk Group 1: Post Office
- Diagnostic Specimens, Risk Group 2,3: Post Office
- Diagnostic Specimens: Transport via Air

## Appendix A

### ICAO GUIDANCE DOCUMENT

#### Consignment of Diagnostic Specimens 2003-2004

Interpretation/Guidance Document developed by ICAO Dangerous Goods Panel members nominated by Canada, United Kingdom and United States in collaboration with the World Health Organization

Note: This document is only valid for the period of 1 January 2003 through 31 December 2004. Please refer to the ICAO website for updates and changes that have occurred since this document was published ( <http://www.icao.org> )

#### Introduction

The 2003-2004 ICAO Technical Instructions include amendments for diagnostic specimens. The purpose of this document is to provide information and guidance for complying with the amendments. Specifically the document provides guidance on:

- Use of the new requirements for diagnostic specimens
- Packaging and consignment procedures
- Passenger and operator provisions
- Substances included or excluded from shipment as diagnostic specimens
- Emergency response procedures

The previous references to risk groups for determining if a substance may be transported as a diagnostic specimen have been removed (see 2;6.3.1.3.2) The 2003-2004 edition of the Technical Instructions maintains the risk group criteria for classifying infectious substances but it is anticipated that the classification criteria will be replaced in the 2005-2006 edition of the Technical Instructions when the ICAO Dangerous Goods Panel considers the infectious substances requirements that were recently adopted for the 13<sup>th</sup> revised edition of the UN Model Regulations. As a result of the 2003-2004 amendments, specimens known or suspected of containing pathogens meeting the criteria for risk groups 2 or 3 may be transported as diagnostic specimens when they are transported for diagnostic or investigational purposes. Specimens known or suspected of containing risk group 4 pathogens must be classified in Division 6.2 under UN 2814 or UN 2900, as appropriate and transported according to the requirements for these substances.

The text below is provided to explain the impact of the amendments to the diagnostic specimens requirements in the Technical Instructions. The new requirements for diagnostic specimens that were adopted by the 12<sup>th</sup> revised edition of the UN Model Regulations have been adopted in other modal regulations and in certain national and regional transport regulations effective January 1, 2003.

#### The definition and relevant requirements

6.3.1.3.1 Diagnostic specimens are any human or animal material including, but not limited to, excreta, secretions, blood and its components, tissue and tissue fluids being transported for diagnostic or investigational purposes, but excluding live infected animals.

6.3.1.3.2 Diagnostic specimens must be assigned to UN 3373 unless the source patient or animal has or may have a serious human or animal disease which can be readily transmitted from one individual to another, directly or indirectly, and for which effective treatment and preventative measures are not usually available, in which case they must be assigned to UN 2814 or UN 2900

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***Note 1.** — Blood which has been collected for the purpose of blood transfusion or for the preparation of blood products, and blood products and any tissues or organs intended for use in transplants are not subject to these Instructions.*

***Note 2.** — Assignment to UN 2814 or UN 2900 must be based on known medical history of the patient or animal, endemic local conditions, symptoms of the patient or animal, or professional judgment concerning individual circumstances of the patient or animal.*

Diagnostic specimens, including those taken from apparently healthy individuals, may contain pathogens that meet the criteria for risk groups 1, 2, 3 or 4. Pathogens are defined as microorganisms (including bacteria, viruses, rickettsiae, parasites, fungi) and other agents such as prions that can cause disease in humans or animals. Pathogens are carried in blood, on the skin, in saliva or feces. Specimens containing risk group 1 pathogens are not subject to the Technical Instructions. Specimens containing risk group 4 pathogens are not permitted for transport as diagnostic specimens. Diagnostic specimens containing risk group 2 or 3 pathogens present a lower risk in transport as compared to infectious substances containing risk group 4 pathogens or pathogens that are intentionally propagated in high concentrations such as those being transported for medical research. Effective treatments are available and the risk of the spread of infection is limited for risk group 2 or 3 pathogens. Additionally, the risk of transmission from one infected individual to another is not as great for these pathogens. Since the packaging requirements of packing instruction 650 afford a high level of safety the probability of exposure is relatively low. The probability of transmission of an infection or disease to an exposed individual from a diagnostic specimen is also relatively low. Effective and cautious emergency response procedures and employee training significantly minimize the risk of exposure and subsequent transmission of infection or disease.

Consignors, who would normally be health care professionals, must make a judgment about the presence of pathogens of risk group 4. However, such judgment is not required in respect of risk group 2 or 3, provided the specimens are being transported for diagnostic or investigational purposes. Specimens containing pathogens of risk group 2 or 3 transported for any other purpose must be consigned as UN2814 or UN2900.

These requirements were developed in coordination with experts from the World Health Organization (WHO) and provide a level of safety commensurate with the risk in transport without imposing an undue burden on those who are required to determine whether an infectious substance may be transported as a diagnostic specimen. In particular the amendments:

- avoid direct reference to WHO Risk Groups, which had been developed by WHO for purposes other than transport and remove ambiguity related to the previous use of the terms “reasonably expected to contain” or “those where a relatively low probability exists” ;
- limit the application of requirements in transport to those commensurate with the actual , rather than the perceived, risk;
- require easily obtainable, suitable packaging affording a high level of safety appropriate to the degree of hazard and conditions of transport. Packing instruction 650 is appropriate for the transport of diagnostic specimens containing pathogens belonging to risk group 2 and 3;

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- permit ready consignment and provide for the universal and effective treatment of individuals in the healthcare system.

It should be noted that determining if a substance is infectious has always included subjective analysis in the absence of actual testing. The 2003-2004 amendment minimizes the subjectivity relative to determining if a substance may be transported as a diagnostic specimen. Classifying these materials based on the level of risk and applying transport requirements commensurate with that risk should ensure an adequate level of safety.

### **Packaging and consignment procedures**

Packing Instruction 650 is intended to provide all the information necessary to prepare and transport safely a consignment of diagnostic specimens. Among other requirements:

1) the packaging must be of good quality capable of passing a 1.2m drop test and must consist of three components:

a primary receptacle containing the diagnostic specimen;

a secondary packaging, and

an outer packaging with suitable cushioning material.

2) Either the primary or secondary receptacle must be capable of withstanding an internal pressure producing a pressure differential of not less than 95kPa for liquids.

3) The package must be marked "DIAGNOSTIC SPECIMEN". The UN number is not required to be shown.

### **Passenger and Operator Provisions**

Diagnostic specimens are not permitted for transport in carry-on or checked baggage and must not be carried on a person. Operators must not load or transport diagnostic specimens unless they are transported as cargo in accordance with the provisions of 7, 2.1 of the Technical Instructions.

### **Substances Excluded From Shipment As Diagnostic Specimens**

**NOTE 1:** *The following list is not exhaustive. Infectious substances, including those containing new or emerging pathogens, which do not appear in the following list but which meet the same criteria must not be transported as a diagnostic specimen. In addition, if there is doubt as to whether or not a pathogen falls within this category it must not be transported as a diagnostic specimen.*

**NOTE 2:** *In the following table, the microorganisms indicated in italics are bacteria, mycoplasmas, rickettsiae or fungi.*

**NOTE 3: Cultures** (laboratory stocks) are the result of a process by which pathogens are amplified or propagated in order to generate high concentrations, thereby increasing the risk of infection when exposure to them occurs. This refers to cultures prepared for the intentional generation of pathogens and does not include cultures intended for diagnostic and clinical purposes. Cultures prepared for the intentional generation of pathogens may not be transported as diagnostic specimens.

**NOTE 4:** *If a health authority list is available that shows other pathogens regarded as Risk Group 4 this should also be taken into account and the substances should not be transported as diagnostic specimens.*

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INDICATIVE EXAMPLES OF INFECTIOUS SUBSTANCES FORBIDDEN AS DIAGNOSTIC SPECIMENS IN ANY FORM UNLESS OTHERWISE INDICATED	
UN Number and Proper Shipping Name	Micro-organism
Infectious substances affecting humans	<i>Bacillus anthracis</i> (cultures only) <i>Brucella abortus</i> (cultures only) <i>Brucella melitensis</i> (cultures only) <i>Brucella suis</i> (cultures only) <i>Burkholderia mallei</i> - <i>Pseudomonas mallei</i> – Glanders (cultures only) <i>Burkholderia pseudomallei</i> – <i>Pseudomonas pseudomallei</i> (cultures only) <i>Chlamydia psittaci</i> - avian strains (cultures only) <b><i>Clostridium botulinum</i> (cultures only)</b> <i>Coccidioides immitis</i> (cultures only) <i>Coxiella burnetii</i> (cultures only) Crimean-Congo hemorrhagic fever virus Dengue virus (cultures only) Eastern equine encephalitis virus (cultures only) <i>Escherichia coli</i> , verotoxigenic (cultures only) Ebola virus Flexal virus <i>Francisella tularensis</i> (cultures only) Guanarito virus Hantaan virus Hantaviruses causing hantavirus pulmonary syndrome Hendra virus Hepatitis B virus (cultures only) Herpes B virus (cultures only) Human immunodeficiency virus (cultures only) Highly pathogenic avian influenza virus (cultures only) Japanese Encephalitis virus (cultures only) Junin virus Kyasanur Forest disease virus Lassa virus Machupo virus Marburg virus Monkeypox virus <i>Mycobacterium tuberculosis</i> (cultures only) Nipah virus Omsk hemorrhagic fever virus Poliovirus (cultures only) Rabies virus <i>Rickettsia prowazekii</i> (cultures only) <i>Rickettsia rickettsii</i> (cultures only) Rift Valley fever virus Russian spring-summer encephalitis virus (cultures only) Sabia virus <i>Shigella dysenteriae</i> type 1 (cultures only) Tick-borne encephalitis virus (cultures only)

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INDICATIVE EXAMPLES OF INFECTIOUS SUBSTANCES FORBIDDEN AS DIAGNOSTIC SPECIMENS IN ANY FORM UNLESS OTHERWISE INDICATED	
UN Number and Proper Shipping Name	Micro-organism
Infectious substances affecting humans ( <i>cont'd</i> )	Variola virus Venezuelan equine encephalitis virus West Nile virus (cultures only) Yellow fever virus (cultures only) <i>Yersinia pestis</i> (cultures only)
2900 Infectious substances affecting animals	African horse sickness virus African swine fever virus Avian paramyxovirus Type 1 - Newcastle disease virus Bluetongue virus Classical swine fever virus Foot and mouth disease virus Lumpy skin disease virus <i>Mycoplasma mycoides</i> - Contagious bovine pleuropneumonia Peste des petits ruminants virus Rinderpest virus Sheep-pox virus Goatpox virus Swine vesicular disease virus Vesicular stomatitis virus

## Emergency Response Procedures

### Mitigation procedures:

Isolate spill or leak area immediately in all directions.

Keep unauthorized personnel away.

Obtain identity of substance involved if possible and report the spill to the appropriate authorities.

Do not touch or walk through spilled material.

Do not touch damaged containers or spilled material unless wearing appropriate protective clothing.

Be particularly careful to avoid contact with broken glass or sharp objects that may cause cuts or abrasions that could significantly increase the risk of exposure.

Damaged packages containing solid CO<sub>2</sub> as a refrigerant may produce water or frost from condensation of air. Do not touch this liquid as it could be contaminated by the contents of the parcel.

Liquid nitrogen may be present and can cause severe burns.

Absorb spilled materials with earth, sand or other non-combustible material while avoiding direct contact.

Cover damaged package or spilled material with damp towel or rag and keep wet with liquid bleach or other disinfectant. Liquid bleach will generally effectively inactivate the released substance.

**DO NOT CLEAN-UP OR DISPOSE OF, EXCEPT UNDER SUPERVISION OF A SPECIALIST.**

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### **First Aid:**

Move exposed person(s) to a safe isolated area.

**CAUTION: Exposed person(s) may be a source of contamination.**

Call emergency medical services.

Remove and isolate contaminated clothing and shoes.

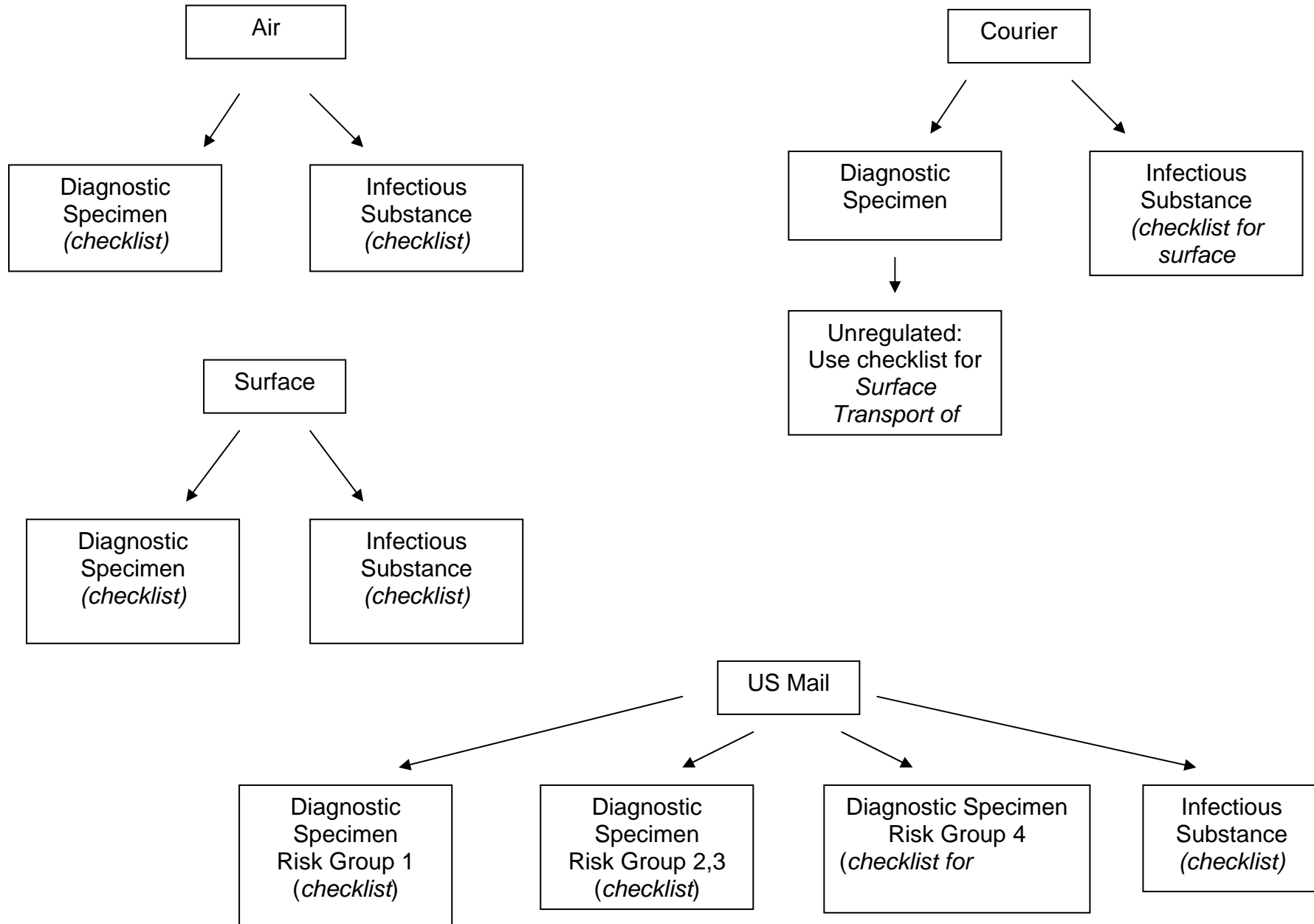
In case of contact with substance, immediately flush skin or eyes with running water for at least 20 minutes.

Effects of exposure (inhalation, ingestion or skin contact) to substance may be delayed.

For further assistance, contact the appropriate public health authority.

Ensure that medical personnel are aware of the substances involved, and take precautions to protect themselves.

Appendix A  
**METHOD OF TRANSPORT**





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<b>INFECTIOUS SUBSTANCE: SURFACE (TAXI, PRIVATE CAR, COURIER)</b>	
<b>Packaging Checklist</b> { Documented Training is required prior to packaging and shipping infectious Agents: 49 CFR 172.700 (h), IATA Section 1.5 }	
49 CFR 173.196	Triple packaging, primary and secondary are leak-proof for liquids and sift-proof for solids (utilize commercially available shipping systems).
49 CFR 173.196 IATA 602	In ambient or higher temperature, primary receptacles have been heat-sealed; have a skirted stopper or a metal crimp seal. Screw caps have been reinforced with adhesive tape.
42 CFR 72.3 (b)	<b>Quantities:</b> a) Max. single primary container or several primary containers 1000 mL. b) Max. inside an outer container 4000 mL.
42 CFR 72.3 (b)	If volume exceeds 50 mL shock absorbent material was inserted at top, bottom and sides between secondary container and outer shipping container.
	Paperwork is separated from the specimen by a plastic sleeve or bag.
49 CFR 173.196	Absorbent material capable of containing an entire spill, was placed between primary and secondary receptacles. Contact between primary receptacles was prevented.
49 CFR 173.196	Multiple primaries placed in secondary packaging must be wrapped individually to prevent contact with each other.
49 CFR 173.196 IATA 602	The primary receptacle or secondary packaging used for infectious substances must be capable of withstanding, without leakage, an internal pressure producing a pressure differential of not less than 95 kPa and at temperatures of –40 to 55 °C (utilize commercially available shipping systems).
49 CFR 178.503	Certified outer shipping package meets UN class 6.2 specifications and packaging instructions PI 602 and bears the UN Packaging Specification Marking. Packaging systems must be 4G Class 6.2 and include the last two digits of the year of manufacture (utilize commercially available shipping systems).
49 CFR 173.196 IATA 602	Outer packaging is at least 100 mm in overall external dimensions.
49 CFR 173.196 IATA 602	An itemized list of contents is enclosed between secondary packaging and outer packaging.
49 CFR 173.199	Interior supports in place to secure secondary package after ice has dissipated or melted (utilize commercially available shipping systems).
49 CFR 173.196	Chemical Ice, dry ice, or wet ice (if applicable) has been placed outside the secondary package (Wet ice should only be used for same day delivery)
49 CFR 173.196	For wet ice, is the package leak-proof? (sealed in plastic bag) For dry ice, does packaging permit release of carbon dioxide gas? (if applicable ) (utilize commercially available shipping systems).
<b>Marking and Labeling Requirements</b>	
49 CFR 172.312	Orientation (Up) arrows on opposite sides of shipping container if primary containers contain greater than 50 mL of liquid.
49 CFR 172.400 Table 49 CFR 172.101	A UN shipping name label: “Infectious substance, affecting humans (organisms’ technical names), UN 2814”
49 CFR 172.432	Diamond shaped Class 6 Infectious Substance label with CDC phone number.

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	(2" x 2" ok for only the smallest size package)
49 CFR 172.446	Dry Ice: Diamond shaped Class 9 label placed on outer packaging. Enter weight in kg.
49 CFR 172.301 (d)	Consignee's or Consignor's name and address required if package will be transferred to another carrier.
DOT/IATA	Overpacks (not to be confused with outer packaging), if used, must have all the labeling of inner packagings plus a label or statement that "inner packagings comply with prescribed specifications".
<b>Shipper's Declaration of Dangerous Goods</b>	
49 CFR 172.301 (d)	Shipper's name and address
49 CFR 172.301 (d)	Consignee's name and address
	Cross out "Radioactive" under shipment type
49 CFR 172.202 Table 172.101	<b>Proper Shipping Name:</b> "Infectious Substance, Affecting Humans (technical name of organisms)" "Dry Ice" (if applicable)
49 CFR 172.202 Table 49 CFR 172.101	<b>Class or Division:</b> "6.2" for organisms "9" for Dry ice (if applicable)
49 CFR 172.301	<b>UN or ID number:</b> "UN2814" for organisms "UN1845" for Dry ice (if applicable)
49 CFR 172.202 (a)(4)	<b>Packing Group:</b> "III" for Dry ice (if applicable)
49 CFR 172.202 Table 49 CFR 172.101	<b>Quantity and type of Packing:</b> e.g., "1 x 50 mL" for organisms e.g. "3 kg" for Dry Ice (if applicable) "Packed in one fiberboard box"
IATA 602 IATA 904	<b>Packing Instructions:</b> Infectious Substance.....602 Dry Ice.....904
49 CFR 172.604 (d)	<b>Additional Handling information:</b> "Emergency Contact: (name) (phone numbers)" {Must be a 24/7 number}
	Name/Title of Signatory:
	Place and Date:
	Signature: (make sure you are in compliance before signing)
<b>Additional</b>	
CAP Requirement	Prior to shipment notify the Washington State Public Health Lab of its arrival time. Email: <a href="mailto:PHL.mailroom@doh.wa.gov">PHL.mailroom@doh.wa.gov</a> Phone: (206) 418-5579 FAX No.: (206) 418-5405
42 CFR 72.3 (f)	You must keep a copy of a receipt of delivery
42 CFR 72.4	You must notify the Director, CDC, if shipment was not received within 5 days
49 CFR 172.201 (e)	You must retain a copy of the shipping paper for 375 days after acceptance by the carrier. It must include the date of acceptance by the carrier..

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<b>INFECTIOUS SUBSTANCE: TRANSPORT VIA AIR</b>	
<b>Packaging Checklist</b> { Documented Training is required prior to packaging and shipping infectious Agents: 49 CFR 172.700 (h), IATA Section 1.5 }	
49 CFR 173.196	Triple packaging; primary and secondary are leak-proof for liquids and sift-proof for solids (utilize commercially available shipping systems).
49 CFR 173.196 IATA 602	In ambient or higher temperature, primary receptacles have been heat-sealed, have a skirted stopper or a metal crimp seal. Screw caps have been reinforced with adhesive tape.
Table 49 CFR 172.101 49 CFR 172.102(c)(1)	<b>Quantities:</b> (unless meet Special provisions A81 or A82) Max. 50 mL or 50 gms for passenger aircraft Max. 500 mL or 500 gms primary and 4 L or 4 kgs for total package for Cargo aircraft
	Paperwork is separated from the specimen by a plastic sleeve or bag.
49 CFR 173.196	Absorbent material capable of containing an entire spill, was placed between primary and secondary receptacles.
49 CFR 173.196	Multiple primaries placed in secondary packaging must be wrapped individually to prevent contact with each other.
49 CFR 173.196 IATA 602	The primary receptacle or secondary packaging used for infectious substances must be capable of withstanding, without leakage, an internal pressure producing a pressure differential of not less than 95 kPa and at temperatures of –40 to 55 °C (utilize commercially available shipping systems).
49 CFR 178.503	Certified outer shipping package meets UN class 6.2 specifications and packaging instructions PI 602 and bears the UN Packaging Specification Marking. Packaging systems must be 4G Class 6.2 and include the last two digits of the year of manufacture (utilize commercially available shipping systems).
49 CFR 173.196 IATA 602	Outer packaging is at least 100 mm in overall external dimensions.
49 CFR 173.196 IATA 602	An itemized list of contents is enclosed between secondary packaging and outer packaging.
49 CFR 173.199	Interior supports in place to secure secondary package after ice has dissipated or melted (utilize commercially available shipping systems).
49 CFR 173.196	Chemical Ice, dry ice, or wet ice (if applicable) has been placed outside the secondary package (Wet ice should only be used for same day delivery)
49 CFR 173.196	For wet ice, is the package leak-proof? (sealed in plastic bag) For dry ice, does packaging permit release of carbon dioxide gas? (if applicable ) (utilize commercially available shipping systems).
<b>Marking and Labeling Requirements</b>	
49 CFR 172.312 IATA 602 Section 7	Orientation (Up) arrows on opposite sides of shipping container if primary containers contain greater than 50 mL of liquid.
49 CFR 172.400, 49 CFR 172.101, IATA 7.1	A UN shipping name label: “Infectious substance, affecting humans (organisms’ technical names), UN 2814” and the volume/weight of the sample.

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49 CFR 172.432	Diamond shaped Class 6 Infectious Substance label with CDC phone number. (2" x 2" ok for only the smallest size package)
IATA	For volumes over 50 mL (and special provisions A81 and A82 are not applicable) Cargo only label (orange danger label) was placed adjacent to Class 6 label. (2" x 2" ok for only the smallest size package)
49 CFR 172.446	Dry Ice: Diamond shaped Class 9 label placed on outer packaging. Enter weight in Kg.
IATA 602	Shipper's name, address and telephone number on box. Consignee's name and address on box.
DOT/IATA	Overpacks (not to be confused with outer packaging), if used, must have all the labeling of inner packagings plus a label or statement that "inner packagings comply with prescribed specifications".
<b>Shipper's Declaration of Dangerous Goods</b> ( <i>Download and type, do not hand write</i> )	
49 CFR 172.301 (d)	Shipper's name and address
49 CFR 172.301 (d)	Consignee's name and address
	Cross out "Radioactive" under shipment type
	Cross out "Passenger aircraft" or "Cargo Aircraft" depending on quantities
49 CFR 172.202 Table 172.101	<b>Proper Shipping Name:</b> "Infectious Substance, Affecting Humans (technical name of organisms)" "Dry Ice" (if applicable)
49 CFR 172.202 Table 49 CFR 172.101	<b>Class or Division:</b> "6.2" for organisms "9" for Dry ice (if applicable)
49 CFR 172.301	<b>UN or ID number:</b> "UN2814" for organisms "UN1845" for Dry ice (if applicable)
49 CFR 172.202 (a)(4)	<b>Packing Group:</b> "III" for Dry ice (if applicable)
49 CFR 172.202 Table 49 CFR 172.101	<b>Quantity and type of Packing:</b> e.g. "1 x 50 mL" for organisms e.g. "3 kg" for Dry Ice (if applicable) "Packed in one fiberboard box"
IATA 602 IATA 904	<b>Packing Instructions:</b> Infectious Substance.....602 Dry Ice.....904
49 CFR 172.604 (d) IATA 602	<b>Additional Handling information:</b> "Emergency Contact: (name) (phone numbers)" {Must be a 24/7 number} "Prior arrangements as required by the IATA Dangerous Goods Reg 1.3.3.1 have been made" {Shipper is required to make advance arrangements with consignee and the carrier to ensure that shipment is transported and delivered without delay}
	Name/Title of Signatory:
	Place and Date:
	Signature: (make sure you are in compliance before signing)

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Additional	
CAP Requirement	Prior to shipment notify the Washington State Public Health Lab of its arrival time. Email: <a href="mailto:PHL.mailroom@doh.wa.gov">PHL.mailroom@doh.wa.gov</a> Phone: (206) 418-5579 FAX No.: (206) 418-5405
42 CFR 72.3 (f)	You must keep a copy of a receipt of delivery
42 CFR 72.4	You must notify the Director, CDC, if shipment was not received within 5 days
49 CFR 172.201 (e)	You must retain a copy of the shipping paper for 375 days after acceptance by the carrier. It must include the date of acceptance (keep the airbill).

**A81:** The quantity limits do not apply to body fluids known to contain or suspected of containing an infectious substance when transported in primary receptacles not exceeding 1000 mL and in outer packagings not exceeding 4 L and packaged according to 49 CFR 173.196.

**A82:** The quantity limits do not apply to human or animal body parts, whole organs or whole bodies known to contain or suspected of containing an infectious substance.

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<b>INFECTIOUS SUBSTANCE: POST OFFICE</b>	
<b>Packaging Checklist</b> <i>{ Documented Training is required prior to packaging and shipping infectious Agents:  49 CFR 172.700 (h), IATA Section 1.5 }</i>	
49 CFR 173.196 Fed Reg. 67 No.244	Triple packaging; primary and secondary are leak-proof for liquids and sift-proof for solids <i>(utilize commercially available shipping systems)</i> .
49 CFR 173.196 IATA 650	In ambient or higher temperature, primary receptacles have been heat-sealed, have a skirted stopper or a metal crimp seal. Screw caps have been reinforced with adhesive tape.
49 CFR 173.6 amendment IATA 650 Fed Reg. 67 No.244	<b>Quantities:</b> Max. 50 mL total per mailpiece.
	Paperwork is separated from the specimen by a plastic sleeve or bag.
49 CFR 173.196 IATA 650 Fed Reg. 67 No.244	Absorbent material capable of containing an entire spill, was placed between primary and secondary receptacles.
49 CFR 173.196	Multiple primaries placed in secondary packaging must be wrapped individually to prevent contact with each other.
49 CFR 173.196 IATA 650 Fed Reg. 67 No.244	The primary receptacle or secondary packaging used for infectious substances must be capable of withstanding, without leakage, an internal pressure producing a pressure differential of not less than 95 kPa and at temperatures of –40 to 55 °C <i>(utilize commercially available shipping systems)</i> .
49 CFR 178.503 49 CFR 178.609 Fed Reg. 67 No.244	Certified outer shipping package meets UN class 6.2 specifications and packaging instructions PI 602 and bears the UN Packaging Specification Marking. Packaging systems must be 4G Class 6.2 and include the last two digits of the year of manufacture <i>(utilize commercially available shipping systems)</i> .
49 CFR 173.196 IATA 650 Fed Reg. 67 No.244	Outer packaging is at least 100 mm in overall external dimensions. Completed package must meet drop test <i>(utilize commercially available shipping systems)</i> .
49 CFR 173.196 IATA 602	An itemized list of contents is enclosed between secondary packaging and outer packaging.
49 CFR 173.199	Interior supports in place to secure secondary package after ice has dissipated or melted <i>(utilize commercially available shipping systems)</i> .
49 CFR 173.196 IATA 602	Chemical Ice, dry ice, or wet ice <i>(if applicable)</i> has been placed outside the secondary package (Wet ice should only be used for same day delivery)  Air Transport: Max. 5 pounds  Surface: no restriction
49 CFR 173.196 IATA 602 Fed Reg. 67 No.244	For wet ice, is the package leak-proof? (sealed in plastic bag) For dry ice, does packaging permit release of carbon dioxide gas? <i>(if applicable ) (utilize commercially available shipping systems)</i>

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Marking and Labeling Requirements	
Fed Reg. 67 No. 244	Biohazard warning labels are attached to secondary packaging.
42 CFR 72.3 (d) Fed Reg. 67 No.244	Etiologic Agents/Biohazard Material label was placed on address side of the mailpiece.
49 CFR 172.312 39 CFR Part 111	Orientation (Up) arrows on opposite sides of shipping container for liquids.
49 CFR 172.400 Fed Reg. 67 No.244	A UN shipping name label: “Infectious substance, affecting humans ( <i>organisms’ technical names</i> ), UN 2814” was placed on the address side of the mailpiece.
49 CFR 172.432	Diamond shaped Class 6 Infectious Substance label with CDC phone number. (2” x 2” <i>ok for only the smallest size package</i> )
49 CFR 172.446 Fed Reg. 67 No.244	Dry Ice: Diamond shaped Class 9 label ( <i>if applicable</i> ). “Dry Ice, UN1845” and its weight was placed on address side of mailpiece.
IATA 602	Shipper’s name and address on box.. Consignee’s name and address on box.
Shipper’s Declaration of Dangerous Goods (USPS C023 1.9)	
49 CFR 172.301 (d)	Shipper’s name and address
49 CFR 172.301 (d)	Consignee’s name and address
	Cross out “Radioactive” under shipment type
	Cross out “Passenger aircraft” or “Cargo Aircraft” depending on quantities
49 CFR 172.202 Table 172.101	<b>Proper Shipping Name:</b> “Infectious Substance, Affecting Humans ( <i>technical name of organisms</i> )” “Dry Ice” ( <i>if applicable</i> )
49 CFR 172.202 Table 49 CFR 172.101	<b>Class or Division:</b> “6.2” for organisms “9” for Dry ice ( <i>if applicable</i> )
49 CFR 172.301	<b>UN or ID number:</b> “UN2814” for organisms ”UN1845” for Dry ice ( <i>if applicable</i> )
49 CFR 172.202 (a)(4)	<b>Packing Group:</b> “III” for Dry ice ( <i>if applicable</i> )
49 CFR 172.202 Table 49 CFR 172.101	<b>Quantity and type of Packing:</b> e.g. “1 x 50 mL” for organisms e.g. “3 kg” for Dry Ice ( <i>if applicable</i> ) “Packed in one fiberboard box”
IATA 602 IATA 904	<b>Packing Instructions:</b> Infectious Substance.....602 Dry Ice.....904
49 CFR 172.604 (d) IATA 602	<b>Additional Handling information:</b> “Emergency Contact: ( <i>name</i> ) ( <i>phone numbers</i> )” { <i>Must be a 24/7 number</i> } “Prior arrangements as required by the IATA Dangerous Goods Reg 1.3.3.1 have

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	been made” { <i>Shipper is required to make advance arrangements with consignee and the carrier to ensure that shipment is transported and delivered without delay</i> }
	Name/Title of Signatory:
	Place and Date:
	Signature: <i>(make sure you are in compliance before signing)</i>
<b>Additional</b>	
CAP Requirement	Prior to shipment notify the Washington State Public Health Lab of its arrival time. Email: <a href="mailto:PHL.mailroom@doh.wa.gov">PHL.mailroom@doh.wa.gov</a> Phone: (206) 418-5579 FAX No.: (206) 418-5405
USPS C023 8.3 USPS C023 1.9	<i>To be sent by Registered mail: First Class or Priority. The shipping papers have been signed in triplicate by the mailer and affixed to the outside of the mailpiece within an envelope that can be easily opened and resealed.</i>
42 CFR 72.3 (f)	A copy of a receipt of delivery has been kept.
49 CFR 172.201 (e)	<i>You must retain a copy of the shipping paper for 375 days after acceptance by the carrier. It must include the date of acceptance by the Post Office.</i>
42 CFR 72.4	You must notify the Director, CDC, if shipment was not received within 5 days



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<b>DIAGNOSTIC SPECIMENS, RISK GROUP 1: POST OFFICE</b>	
<b>Packaging Checklist</b>	
{ Documented Training is required prior to packaging and shipping infectious Agents: 49 CFR 172.700 (h), IATA Section 1.5 }	
49 CFR 173.199 (a)(1)	Triple packaging; primary and secondary is leak-proof for liquids and sift-proof for solids (utilize commercially available shipping systems).
49 CFR 173.6 amendment	<b>Quantities:</b> Max. each inner package 0.5 kg or 0.5 L and Max. outer packaging 4 kg or 4 L Max. single inner package 16 kg or 16 L, in a single outer packaging.
	Paperwork is separated from the specimen by a plastic sleeve or bag.
49 CFR 173.196	Absorbent material capable of containing an entire spill, was placed between primary and secondary receptacles. Contact between primary receptacles was prevented.
49 CFR 173.196	An itemized list of contents is enclosed between secondary packaging and outer packaging.
49 CFR 173.199	Interior supports in place to secure secondary package after ice has dissipated or melted (utilize commercially available shipping systems).
49 CFR 178.603	Completed package passes drop test of 1.2 meters.
49 CFR 173.196	Chemical Ice, dry ice, or wet ice (if applicable) has been placed outside the secondary package (Wet ice should only be used for same day delivery)
49 CFR 173.196	For wet ice, is the package leak-proof? (sealed in plastic bag) For dry ice, does packaging permit release of carbon dioxide gas? (if applicable) (utilize commercially available shipping systems).
<b>Marking and Labeling Requirements</b>	
OSHA: 1910.103 0(g)(1)(i)(A)	Biohazard warning label attached to secondary packaging (not outside box).
49 CFR 172.312 IATA 602 Section 7	Orientation (Up) arrows on opposite sides of shipping container if primary containers contain greater than 50 mL of liquid.
49 CFR	Outer package marked with "Diagnostic Specimen".
49 CFR 172.446	Dry Ice: Diamond shaped Class 9 label placed on outer packaging. Enter weight in Kg.
49 CFR 172.301 (d) IATA 650	Shipper's name, address and telephone number on box. Consignee's name and address on box.
DOT/IATA	Overpacks (not to be confused with outer packaging), if used, has all the labeling of inner packagings plus this label "Inner packagings comply with prescribed specifications".
<b>Documentation</b>	
CAP Requirement	Prior to shipment notify the Washington State Public Health Lab of its arrival time. Email: <a href="mailto:PHL.mailroom@doh.wa.gov">PHL.mailroom@doh.wa.gov</a> Phone: (206) 418-5579 FAX No.: (206) 418-5405
49 CFR 172.201 (e)	You must retain a copy of the shipping paper for 375 days after acceptance by the carrier. It must include the date of acceptance by the carrier.

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### Packaging Checklist

{ Documented Training is required prior to packaging and shipping infectious Agents:  
49 CFR 172.700 (h), IATA Section 1.5 }

Fed Reg. 67 No. 244	Double packaging for 50 mL or less with a securely sealed primary receptacle. The secondary container is leakproof and may serve as the outer shipping container. Triple packaging when exceeding 50 mL with max. 1000 mL per primary and secondary container. Secondary container cannot serve as outer shipping container. Max. per mailpiece 4,000 mL.
	Paperwork is separated from the specimen by a plastic sleeve or bag.
Fed Reg. 67 No. 244	Absorbent material capable of containing an entire spill and cushioning material to protect from breakage surrounds the primary receptacle.
<b>Marking and Labeling Requirements</b>	
Fed Reg. 67 No. 244	Biohazard warning label attached to each secondary container.
<b>Documentation</b>	
CAP Requirement	Prior to shipment notify the Washington State Public Health Lab of its arrival time. Email: <a href="mailto:PHL.mailroom@doh.wa.gov">PHL.mailroom@doh.wa.gov</a> Phone: (206) 418-5579 FAX No.: (206) 418-5405
Fed Reg. 67 No. 244	To be sent First Class or Priority Mail or Package services.

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### DIAGNOSTIC SPECIMENS, RISK GROUP 2,3: POST OFFICE

#### Packaging Checklist

*{Documented Training is required prior to packaging and shipping infectious Agents:  
49 CFR 172.700 (h), IATA Section 1.5}*

49 CFR 173.196 Fed Reg. 67 No. 244	Triple packaging; primary and secondary are leak-proof for liquids and sift-proof for solids <i>(utilize commercially available shipping systems)</i> .
49 CFR 173.196 IATA 650	In ambient or higher temperature, primary receptacles have been heat-sealed, have a skirted stopper or a metal crimp seal. Screw caps have been reinforced with adhesive tape.
49 CFR 173.6 amendment IATA 650	<b>Quantities:</b> Max. each inner package 0.5 kg or 0.5 L and Max. outer packaging 4 kg or 4 L.
	Paperwork is separated from the specimen by a plastic sleeve or bag.
49 CFR 173.196 IATA 650	Absorbent material capable of containing an entire spill, was placed between primary and secondary receptacles.
49 CFR 173.196 IATA 650	Multiple primaries placed in secondary packaging must be wrapped individually to prevent contact with each other.
49 CFR 173.196 IATA 650 Fed Reg. 67 No. 244	The primary receptacle or secondary packaging used for infectious substances must be capable of withstanding, without leakage, an internal pressure producing a pressure differential of not less than 95 kPa and at temperatures of –40 to 55 °C <i>(utilize commercially available shipping systems)</i> .
49 CFR 173.196 IATA 650	Outer packaging is at least 100 mm in overall external dimensions. Completed package must meet drop test <i>(utilize commercially available shipping systems)</i> .
49 CFR 173.196 IATA 650	An itemized list of contents is enclosed between secondary packaging and outer packaging.
49 CFR 173.199	Interior supports in place to secure secondary package after ice has dissipated or melted <i>(utilize commercially available shipping systems)</i> .
49 CFR 173.196	Chemical Ice, dry ice, or wet ice <i>(if applicable)</i> has been placed outside the secondary package (Wet ice should only be used for same day delivery)
49 CFR 173.196	For wet ice, is the package leak-proof? (sealed in plastic bag) For dry ice, does packaging permit release of carbon dioxide gas? <i>(if applicable)</i> <i>(utilize commercially available shipping systems)</i> .

#### Marking and Labeling Requirements

OSHA: 1910.103 0(g)(1)(i)(A)	Biohazard warning label attached to each secondary container.
49 CFR 172.312 IATA 602 Section 7	Orientation (Up) arrows on opposite sides of shipping container if primary containers contain greater than 50 mL of liquid.
IATA 650 Fed Reg. 67 No. 244	Address side of outer packaging is marked “Diagnostic Specimens”.
49 CFR 172.446 DMM C023	Dry Ice: Diamond shaped Class 9 label <i>(if applicable)</i> . Enter weight in Kg. If shipping by air via USPS you will need a Shipper’s declaration of Dangerous Goods just for the Dry Ice.

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IATA 602	Shipper's name, address and telephone number on box. Consignee's name and address on box.
DOT/IATA	Overpacks ( <i>not to be confused with outer packaging</i> ), if used, must have all the labeling of inner packagings plus a label or statement that "inner packagings comply with prescribed specifications".
<b>Documentation</b>	
CAP Requirement	Prior to shipment notify the Washington State Public Health Lab of its arrival time. Email: <a href="mailto:PHL.mailroom@doh.wa.gov">PHL.mailroom@doh.wa.gov</a> Phone: (206) 418-5579 FAX No.: (206) 418-5405
USPS C023 8.6	<i>To be sent First Class or Priority.</i> (With Dry Ice a Shipper's Declaration of Dangerous Goods is required.)
<b>Packaging Checklist</b> <i>{ Documented Training is required prior to packaging and shipping infectious Agents: 49 CFR 172.700 (h), IATA Section 1.5 }</i>	
49 CFR 173.196	Triple packaging; primary and secondary are leak-proof for liquids and sift-proof for solids ( <i>utilize commercially available shipping systems</i> ).
49 CFR 173.196 IATA 650	In ambient or higher temperature, primary receptacles have been heat-sealed, have a skirted stopper or a metal crimp seal. Screw caps have been reinforced with adhesive tape.
49 CFR 173.6 amendment IATA 650	<b>Quantities:</b> ( <i>unless meet Special provisions A82</i> ) Max. each inner package 0.5 kg or 0.5 L and Max. outer packaging 4 kg or 4 L, Passenger or Cargo aircraft acceptable.
	Paperwork is separated from the specimen by a plastic sleeve or bag.
49 CFR 173.196 IATA 650	Absorbent material capable of containing an entire spill, was placed between primary and secondary receptacles.
49 CFR 173.196 IATA 650	Multiple primaries placed in secondary packaging must be wrapped individually to prevent contact with each other.
49 CFR 173.196 IATA 650	The primary receptacle or secondary packaging used for infectious substances must be capable of withstanding, without leakage, an internal pressure producing a pressure differential of not less than 95 kPa and at temperatures of –40 to 55 °C ( <i>utilize commercially available shipping systems</i> ).
49 CFR 173.196 IATA 650	Outer packaging is at least 100 mm in overall external dimensions. Completed package must meet drop test ( <i>utilize commercially available shipping systems</i> ).
49 CFR 173.196 IATA 650	An itemized list of contents is enclosed between secondary packaging and outer packaging.
49 CFR 173.199	Interior supports in place to secure secondary package after ice has dissipated or melted ( <i>utilize commercially available shipping systems</i> ).
49 CFR 173.196	Chemical Ice, dry ice, or wet ice ( <i>if applicable</i> ) has been placed outside the secondary package ( <i>Wet ice should only be used for same day delivery</i> )

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49 CFR 173.196	For wet ice, is the package leak-proof? ( <i>sealed in plastic bag</i> ) For dry ice, does packaging permit release of carbon dioxide gas? ( <i>if applicable</i> ) ( <i>utilize commercially available shipping systems</i> ).
<b>Marking and Labeling Requirements</b>	
OSHA: 1910.103 0(g)(1)(i)(A)	Biohazard warning label attached to secondary packaging (not outside box).
49 CFR 172.312 IATA 602 Section 7	Orientation (Up) arrows on opposite sides of shipping container if primary containers contain greater than 50 mL of liquid.
IATA 650	Outer packaging is marked "Diagnostic Specimens"
49 CFR 172.446	Dry Ice: Diamond shaped Class 9 label placed on outer packaging. Enter weight in Kg.
IATA 602	Name and telephone number of person responsible for shipment.
DOT/IATA	Overpacks ( <i>not to be confused with outer packaging</i> ), if used, must have all the labeling of inner packagings plus a label or statement that "inner packagings comply with prescribed specifications".
<b>Documentation</b>	
CAP Requirement	Prior to shipment notify the Washington State Public Health Lab of its arrival time. Email: <a href="mailto:PHL.mailroom@doh.wa.gov">PHL.mailroom@doh.wa.gov</a> Phone: (206) 418-5579 FAX No.: (206) 418-5405
IATA 650 and 904	<b>Airbill:</b> In the Nature and Quantity of Goods box place, "Diagnostic Specimen packed in Compliance with IATA Packing Instruction 650" and if applicable: "Dry Ice".
49 CFR 172.201 (e)	<i>You must retain a copy of the shipping paper for 375 days after acceptance by the carrier. It must include the date of acceptance (keep the airbill).</i>
42 CFR 72.3 (f)	You must keep a copy of a receipt of delivery

**A82:** The quantity limits do not apply to human or animal body parts, whole organs or whole bodies known to contain or suspected of containing an infectious substance.

<b>APPENDIX B: PHL Accreditation/Certification</b>	
<b>Accreditation Body</b>	<b>Certification Number</b>
Clinical Laboratory Improvement Act (CLIA)	50D0661453
College Of American Pathologists (CAP)	24626-01
Department Of Energy - Radiation Measurement Laboratory	WN-L074-1
Environmental Protection Agency (EPA) for drinking water bacteriology and environmental/radiation chemistry	WA 00003
Food And Drug Administration (FDA)	FOOD #475 SHELLFISH #705
Medical Test Site License (MTS) WA DOH HSQA Office Of Laboratory Quality Assurance (LQA)	MTS-1327

## Appendix C: Notifiable Conditions

### *Notifiable Conditions & Washington's Laboratories*



The following laboratory results (preliminary or confirmed) are notifiable to public health authorities in Washington in accordance with WAC 246-101. Information provided must include: Specimen Type; Name and Telephone Number of Laboratory; Date Specimen Collected; Date Specimen Received; Requesting Health Care Provider's Name & Telephone Number or Address; Test Result; Name of Patient (if available) or patient identifier; Sex & Date of Birth or Age of Patient (if available).

Blood Lead Level (Elevated) <sup>2&1</sup>  
Blood Lead Level (Non-elevated) <sup>M &1</sup>  
Bordetella pertussis <sup>2\*</sup>  
Brucella <sup>2\*!</sup>  
CD4+ counts <200 or 14% <sup>M &11</sup>  
Chlamydia trachomatis <sup>2\*</sup>  
Clostridium botulinum I <sup>\*!</sup>  
Corynebacterium diphtheriae <sup>2\*!</sup>  
Cryptosporidium parvum <sup>2\*</sup>  
Cyclospora cayentanensis <sup>2\*!</sup>  
Diseases of Suspected  
  Bioterrorism Origin <sup>1\*!</sup>  
    Bacillus anthracis  
    Orthopoxvirus  
Escherichia coli (Shiga-like toxins only) <sup>2\*!</sup>  
Francisella tularensis <sup>!</sup>  
Hepatitis A (Hepatovirus) <sup>2\*</sup>

Human Immunodeficiency Virus <sup>2 &11</sup>  
  (Western Blot, P-24 Antigen, or viral culture)  
Human Immunodeficiency Virus <sup>M &11</sup>  
  (RNA or DNA Nucleic Acid Tests)  
Listeria <sup>2\*</sup>  
Mycobacterium tuberculosis <sup>2 &111 !@</sup>  
Neisseria gonorrhoeae <sup>2\*</sup>  
Neisseria meningitidis <sup>2\*!</sup>  
Rabies <sup>1\*</sup>  
Rubeola <sup>1\*!</sup>  
Salmonella <sup>2\*!</sup>  
Shigella <sup>2\*!</sup>  
Treponema pallidum <sup>!</sup>  
Unusual Diseases of Public  
  Health Significance <sup>1\*</sup>  
Vibrio cholerae <sup>1\*!</sup>  
Yersinia pestis <sup>1\*!</sup>

#### CODE LEGEND

- <sup>1</sup> Immediately Notifiable
- <sup>2</sup> Notifiable within 2 Work Days
- <sup>M</sup> Notifiable on a Monthly Basis
- <sup>\*</sup>Notifiable to the local health  
  department of the patient's residence
- <sup>&1</sup> Notifiable to DOH Lead Program (360-236-4260)
- <sup>&11</sup> Notifiable to DOH – IDRH Assessment  
  (360-236-3412)
- <sup>&111</sup> Notifiable to DOH – TB Services (360-236-3473)
- <sup>!</sup> Specimen submission required
- <sup>@</sup> Antibiotic Sensitivity Testing (First isolates only)

To report a Notifiable Condition, contact the local health jurisdiction of the patients' residence (see LHJ – Chart), unless the condition is reportable directly to DOH. If the patient's local health jurisdiction is unknown, please notify the local health jurisdiction of the health care provider that ordered the diagnostic test.

If no one is available at the local health jurisdiction and a condition is Immediately Notifiable, please call  
(877) 539-4344

For more information please see WAC 246-101 or see  
[www.doh.wa.gov/nc.htm](http://www.doh.wa.gov/nc.htm)

## Appendix C

# Notifiable Conditions & Washington's Hospitals



The following diagnoses are notifiable to local health authorities in Washington in accordance with WAC 246-101. Timeframes for notification are indicated in footnotes. **Immediately notifiable conditions are indicated in bold** and should be reported when suspected or confirmed. These notifications are for conditions that occur or are treated in the hospital. Hospital laboratories should use the *Notifiable Conditions and Washington's Laboratories* Poster.

Acquired Immunodeficiency Syndrome (AIDS) <sup>3</sup> (including AIDS in persons previously reported with HIV infection)  
**Animal Bites** <sup>1</sup>  
**Botulism** <sup>1</sup>  
 (foodborne, wound, and infant)  
**Brucellosis** <sup>1</sup>  
 Campylobacteriosis <sup>3</sup>  
 Chancroid <sup>3</sup>  
*Chlamydia trachomatis* <sup>3</sup>  
**Cholera** <sup>1</sup>  
 Cryptosporidiosis <sup>3</sup>  
 Cyclosporiasis <sup>3</sup>  
**Diphtheria** <sup>1</sup>  
**Disease of Suspected Bioterrorism Origin (including)** <sup>1</sup>  
 Anthrax  
 Smallpox  
**Disease of Suspected Foodborne Origin** <sup>1</sup> (clusters only)  
**Disease of Suspected Waterborne Origin** <sup>1</sup> (clusters only)  
 Encephalitis, viral <sup>3 1</sup>  
**Enterohemorrhagic *E. coli* including *E. coli* 0157:H7 infection** <sup>1</sup>  
 Giardiasis <sup>3</sup>  
 Gonorrhea <sup>3</sup>  
 Granuloma inguinale <sup>3</sup>  
***Haemophilus influenzae* invasive disease** <sup>1</sup>  
 (under age five, excluding otitis media)  
 Hantavirus Pulmonary Syndrome <sup>3</sup>  
**Hemolytic Uremic Syndrome** <sup>1</sup>  
**Hepatitis A - acute** <sup>1</sup>  
 Hepatitis B - acute <sup>3</sup>; chronic <sup>M</sup> (initial diagnosis only)  
 Hepatitis B - surface antigen + pregnant women <sup>3</sup>  
 Hepatitis C - acute and chronic <sup>M</sup> (initial diagnosis only)  
**Hepatitis, unspecified (infectious)** <sup>1</sup>  
 HIV infection <sup>3</sup>  
 Immunization reactions, severe, adverse <sup>3</sup>  
 Legionellosis <sup>3</sup>

Leptospirosis <sup>3</sup>  
**Listeriosis** <sup>1</sup>  
 Lyme disease <sup>3</sup>  
 Lymphogranuloma venereum <sup>3</sup>  
 Malaria <sup>3</sup>  
**Measles (rubeola)** <sup>1</sup>  
**Meningococcal disease** <sup>1</sup>  
 Mumps <sup>3</sup>  
**Paralytic shellfish poisoning** <sup>1</sup>  
**Pertussis** <sup>1</sup>  
**Plague** <sup>1</sup>  
**Poliomyelitis** <sup>1</sup>  
 Psittacosis <sup>3</sup>  
 Q Fever <sup>3</sup>  
**Rabies** <sup>1</sup>  
 Rabies post-exposure prophylaxis <sup>3</sup>  
**Relapsing fever (*borreliosis*) I**  
**Rubella, (including congenital)** <sup>1</sup>  
**Salmonellosis** <sup>1</sup>  
**Shigellosis** <sup>1</sup>  
 Streptococcus Group A, invasive disease <sup>3</sup>  
 Syphilis <sup>3</sup> (including congenital)  
 Tetanus <sup>3</sup>  
 Trichinosis <sup>3</sup>  
**Tuberculosis** <sup>1</sup>  
 Tularemia <sup>3</sup>  
**Typhus** <sup>1</sup>  
 Vibriosis <sup>3</sup>  
**Yellow Fever** <sup>1</sup>  
 Yersiniosis <sup>3</sup>

**Outbreaks of Chickenpox, Influenzae, Nosocomial Infection and Environmentally Related Disease** <sup>1</sup>  
**Unexplained Critical Illness or Death** <sup>1</sup>  
**Rare Diseases of Public Health Significance** <sup>1</sup>

The following diagnoses are notifiable to the Washington State Department of Health in accordance with WAC 246-101. Timeframes for notification are indicated in footnotes. Immediately notifiable conditions are indicated in **bold** and should be reported when suspected or confirmed.

Asthma, occupational (suspected or confirmed)<sup>M</sup>  
 Call 1-888-66-SHARP  
 Birth Defects – Abdominal Wall Defects, Autism, Cerebral Palsy, Down Syndrome, Hypospadias, Limb Reductions, Neural Tube Defects, Oral Clefts <sup>M</sup> Call (360) 236-3492  
 Gunshot Wounds <sup>M</sup> Call (360) 236-3616  
**Pesticide poisoning (hospitalized, fatal, or cluster)** <sup>1</sup>  
 Call 1-800-732-6985

Notification time frame: <sup>1</sup> **immediately**,  
<sup>3</sup> Within 3 work days, <sup>M</sup> Within one month

## INSERT LOCAL REPORTING INFORMATION HERE

**If no one is available at the local health jurisdiction and a condition is Immediately Notifiable, please call (877) 539-4344**

For more information please see WAC 246-101 or see [www.doh.wa.gov/nc.htm](http://www.doh.wa.gov/nc.htm)



## Appendix C

# Notifiable Conditions & The Health Care Provider



The following diagnoses are notifiable to local health authorities in Washington in accordance with WAC 246-101. Timeframes for notification are indicated in footnotes. **Immediately notifiable conditions are indicated in bold** and should be reported when suspected or confirmed.

Acquired Immunodeficiency Syndrome (AIDS) <sup>3</sup> (including  
*AIDS in persons previously reported with HIV infection*)

**Animal Bites** <sup>1</sup>

**Botulism** <sup>1</sup> (foodborne, wound, and infant)

**Brucellosis** <sup>1</sup>

Campylobacteriosis <sup>3</sup>

Chancroid <sup>3</sup>

*Chlamydia trachomatis* <sup>3</sup>

**Cholera** <sup>1</sup>

Cryptosporidiosis <sup>3</sup>

Cyclosporiasis <sup>3</sup>

**Diphtheria** <sup>1</sup>

**Disease of Suspected Bioterrorism Origin (including)** <sup>1</sup>

**Anthrax**

**Smallpox**

**Disease of Suspected Foodborne Origin** <sup>1</sup> (clusters only)

**Disease of Suspected Waterborne Origin** <sup>1</sup> (clusters only)

Encephalitis, viral <sup>3,1</sup>

**Enterohemorrhagic *E. coli* including *E. coli* 0157:H7**  
**infection** <sup>1</sup>

Giardiasis <sup>3</sup>

Gonorrhea <sup>3</sup>

Granuloma inguinale <sup>3</sup>

***Haemophilus influenzae* invasive disease** <sup>1</sup>  
(under age five, excluding otitis media)

Hantavirus Pulmonary Syndrome <sup>3</sup>

**Hemolytic Uremic Syndrome** <sup>1</sup>

**Hepatitis A - acute** <sup>1</sup>

Hepatitis B - acute <sup>3</sup>; chronic <sup>M</sup> (initial diagnosis only)

Hepatitis B - surface antigen + pregnant women <sup>3</sup>

Hepatitis C - acute and chronic <sup>M</sup> (initial diagnosis only)

**Hepatitis, unspecified (infectious)** <sup>1</sup>

Herpes simplex, genital and neonatal <sup>3</sup> (initial infection only)

HIV infection <sup>3</sup>

Immunization reactions, severe, adverse <sup>3</sup>

Legionellosis <sup>3</sup>

Leptospirosis <sup>3</sup>

**Listeriosis** <sup>1</sup>

Lyme disease <sup>3</sup>

Lymphogranuloma venereum <sup>3</sup>

Malaria <sup>3</sup>

**Measles (rubeola)** <sup>1</sup>

**Meningococcal disease** <sup>1</sup>

Mumps <sup>3</sup>

**Paralytic shellfish poisoning** <sup>1</sup>

**Pertussis** <sup>1</sup>

**Plague** <sup>1</sup>

**Poliomyelitis** <sup>1</sup>

Psittacosis <sup>3</sup>

Q Fever <sup>3</sup>

**Rabies** <sup>1</sup>

Rabies post-exposure prophylaxis <sup>3</sup>

**Relapsing fever (*borreliosis*)** <sup>1</sup>

**Rubella, (including congenital)** <sup>1</sup>

**Salmonellosis** <sup>1</sup>

**Shigellosis** <sup>1</sup>

Streptococcus Group A, invasive disease <sup>3</sup>

Syphilis <sup>3</sup> (including congenital)

Tetanus <sup>3</sup>

Trichinosis <sup>3</sup>

**Tuberculosis** <sup>1</sup>

Tularemia <sup>3</sup>

**Typhus** <sup>1</sup>

Vibriosis <sup>3</sup>

**Yellow Fever** <sup>1</sup>

Yersiniosis <sup>3</sup>

**Unexplained Critical Illness or Death** <sup>1</sup>

**Rare Diseases of Public Health Significance** <sup>1</sup>

The following diagnoses are notifiable to the Washington State Department of Health in accordance with WAC 246-101. Timeframes for notification are indicated in footnotes. Immediately notifiable conditions are indicated in **bold** and should be reported when suspected or confirmed.

Asthma, occupational (suspected or confirmed)<sup>M</sup>  
*Call 1-888-66-SHARP*

Birth Defects – Autism<sup>M</sup> *Call (360) 236-3492*

Birth Defects - Cerebral Palsy<sup>M</sup> *Call (360) 236-3492*

Birth Defects – Fetal Alcohol Syndrome/Fetal Alcohol Effects<sup>M</sup>  
*Call (360) 236-3492*

**Pesticide Poisoning (hospitalized, fatal, or cluster)**<sup>1</sup>  
*Call 1-800-732-6985*

Pesticide Poisoning (other)<sup>3</sup> *Call 1-800-732-6985*

Notification time frame: <sup>1</sup> **Immediately**,  
<sup>3</sup> Within 3 work days, <sup>M</sup> Within one month

## INSERT LOCAL REPORTING INFORMATION HERE

**If no one is available at the local health  
jurisdiction and a condition is Immediately  
Notifiable, please call  
(877) 539-4344**

For more information please see WAC 246-101 or see [www.doh.wa.gov/nc.htm](http://www.doh.wa.gov/nc.htm)

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